

AMERICAN VENOUS FORUM

20th ANNUAL MEETING February 20-23, 2008

Charleston Place, Charleston, South Carolina

AMERICAN VENOUS FORUM COMMITTEES EXECUTIVE COMMITTEE

Seattle Washington Cincinnati, Ohio Springfield, Illinois Evanston, Illinois Newark, New Jersey Potomac, Maryland Indianapolis, Indiana Helsingborg, Sweden Ann Arbor, Michigan Newark, New Jersey Birmingham, Alabama Ann Arbor, Michigan

AMERICAN VENOUS FORUM COMMITTEES 2007-2008

PROGRAM COMMITTEE

Joseph D. Raffetto, M.D. (2008), Chair Lowell S. Kabnick, MD (2009) David L. Gillespie, M.D. (2010) Peter J. Pappas, M.D. (2008), Ex-Officio Robert B. McLafferty, M.D. (2007), Ex-Officio

LOCAL ARRANGEMENTS COMMITTEE

William A. Marston, M.D. (2008), Chair

INTERNATIONAL RELATIONS COMMITTEE

Shunichi Hoshino, M.D. (2007), Chair Kevin G. Burnand, M.D. (2008) Paulo Zamboni, MD (2009) Joann M. Lohr, M.D. (2007), Ex-Officio

MEMBERSHIP COMMITTEE

Audra A. Duncan, M.D. (2008), Chair Elna M. Masuda, MD (2009) Mark D. lafrati, M.D. (2010) Robert B. McLafferty, M.D. (2010), Ex-Officio

HONORARY MEMBERSHIP COMMITTEE

Bo G. Eklof, M.D. (2008), Chair Thomas W. Wakefield, MD (2009) Michael C. Dalsing, M.D. (2010)

COMMITTEE ON ISSUES

Paul R. Cordts, M.D. (2008), Chair Steven Elias, M.D. (2009) Byung B. Lee, MD (2010) Audra A. Duncan, M.D. (2011) Robert B. McLafferty, M.D. (2010), Ex-Officio

AMERICAN VENOUS FORUM COMMITTEES 2007-2008

RESEARCH COMMITTEE

Brajesh K. Lal, M.D. (2008), Chair Joseph D. Raffetto, M.D. (2009) Peter K. Henke, M.D. (2010) William A. Marston (2011) Michael Ricci, M.D. (2012) Robert B. McLafferty (2010), Ex-Officio

NOMINATING COMMITTEE

Bo G. Ekof, M.D. (2008), Chair Thomas W. Wakefield, MD (2009) Michael C. Dalsing, M.D. (2010)

NATIONAL SCREENING COMMITTEE

Mark A. Passman, M.D. (2008), Chair Robert B. McLafferty, M.D. (2008) Thomas W. Rooke, M.D. (2008) Joseph A. Caprini, M.D. (2008) Mark D. lafrati, M.D. (2008) William T. Bohannon, M.D. (2008)

OUTCOMES COMMITTEE

Fedor Lurie, M.D. (2008), Co-Chair Mark H. Meissner, M.D. (2008), Co-Chair Gregory L. Moneta, M.D. (2008) Robert L. Kistner, M.D. (2008) Peter N. Neglen, M.D. (2008) Frank T. Padberg, Jr., M.D. (2008) Robert B. McLafferty, M.D. (2008) Mary C. Proctor, M.D. (2008)

BY-LAWS COMMITTEE

Robert B. McLafferty, M.D. (2010), Co-Chair Ashraf M. Mansour, M.D. (2008), Co-Chair Harold Welch, M.D. (2009)

INDUSTRIAL ADVISORY COMMITTEE

Sandra Shaw, Chair

THE AMERICAN VENOUS FORUM FOUNDATION

The American Venous Forum Foundation was organized in 1988 to support the charitable, educational and scientific purposes of the American Venous Forum.

The Foundation provides the **BSN Jobst Fellowship Award**, the **Sigvaris Traveling Fellowship Award**, the **Servier Fellowship Award** and other significant educational grants to stimulate and recognize excellence in published writing on laboratory and clinical research in the study of venous diseases.

The Foundation also oversees the education and objectives of the Venous Education Institute of North America (VEIN). www.venous-info.org

AMERICAN VENOUS FORUM FOUNDING MEMBERS

Robert W. Barnes, M.D. John J. Cranley, M.D. Ralph G. DePalma, M.D. Lazar J. Greenfield, M.D. Michael Hume, M.D. Robert L. Kistner, M.D. Seshadri Raju, M.D. Charles G. Rob, M.D. D. Eugene Strandness, Jr., M.D. J. Leonel Villavicencio, M.D. John J. Bergan, M.D. W. Andrew Dale, M.D. James A. DeWeese, M.D. Robert W. Hobson, II, M.D. George Johnson, Jr., M.D. John M. Porter, M.D. Norman M. Rich, M.D. Joseph G. Sladen, M.D. David S. Sumner, M.D. James S.T. Yao, M.D.

AVF FOUNDATION BOARD OF DIRECTORS

President	Bo G. Eklof, M.D. (2008)	Helsingborg, Sweden
Vice-President	Thomas Wakefield, M.D. (2008)	Ann Arbor, Michigan
Secretary	Robert B. McLafferty, M.D. (2010)	Springfield, Illinois
Treasurer	Joseph A. Caprini, M.D. (2010)	Evanston, Illinois
Directors	Fedor Lurie, M.D.(2008)	Honolulu, Hawaii
	Peter N. Neglen, M.D. (2008)	Flowood, Mississippi
	John Blebea, M.D. (2008)	Philadelphia, Pennsylvania
Ex-Officio	Michael C. Dalsing, M.D. (2009)	Indianapolis, Indiana

THE AMERICAN VENOUS FORUM WAS ORGANIZED IN COOPERATION WITH MEMBERS OF:

The Society for Vascular Surgery American Association of Vascular Surgery The Canadian Society for Vascular Surgery

WITH THE SUPPORT OF MEMBERS OF

The International Union of Phleboloav The North American Society of Phlebology The Phlebology Society of America Austrian Society for Angiology Benelux Society of Phlebology (Belgium, Netherlands and Luxembourg) European Chapter of The International Society for Cardiovascular Surgery German Society of Phlebology and Proctology Latin American Chapter of The International Society for Cardiovascular Surgery Swiss Society for Phlebology Sociedad Mexicana de Angiologia College Francais de Pathologie Société Francaise de Phlebologie Société Francaise d'Angéiologie Societa Italiana de Patologia Vascolare Pan American Society of Phlebology and Lymphology Sociedad Argentina de Flebologia y Linfologia The Australian and New Zealand Society of Phlebology

THE AMERICAN VENOUS FORUM ANNUAL MEETINGS/PAST PRESIDENTS

Feb. 22-24	New Orleans, LA	John J. Bergan, M.D.
Feb. 21-23		Norman M. Rich, M.D.
Feb. 20-22		Lazar J. Greenfield, M.D.
Feb. 26-28		Michael Hume, M.D.
Feb. 24-26	Orlando, FL	George Johnson, Jr., M.D.
	Hilton Walt Disney World Village	
Feb. 23-25	Maui, Hl	James A. DeWeese, M.D.
	Maui Inter-Continental Resort	
Feb. 23-25	Fort Lauderdale, FL	Robert Hobson, M.D.
	Marriott Harbor Beach	
Feb. 22-24	San Diego, CA	Robert L. Kistner, M.D.
	•	
Feb. 20-23		James S. T. Yao, M.D.
Feb. 19-21		
Feb. 18-21	,	
Feb. 3-6	•	David S. Sumner, M.D.
Feb. 22-25		Anthony J. Comerota, M.D.
	•	
Feb. 21-24		Greaory L. Moneta, M.D.
Feb. 20-23		Peter Gloviczki, M.D.
Feb. 26-29		Frank T. Padbera, M.D.
10012027		
Feb 9-13	,	Bo G Eklöf M D
	•	
Feb 22-26		Thomas W. Wakefield, M.D.
10012220		
Feb 11-17		Michael C. Dalsing, M.D.
	Rancho Bernardo Inn	
	Feb. 22-24 Feb. 21-23 Feb. 20-22 Feb. 23-25 Feb. 23-25 Feb. 20-23 Feb. 19-21 Feb. 3-6 Feb. 22-25 Feb. 21-24 Feb. 3-6 Feb. 22-25 Feb. 21-24 Feb. 22-25 Feb. 20-23 Feb. 21-3 Feb. 22-26 Feb. 14-17	Fairmont HotelFeb. 21-23Coronado, CA

AMERICAN VENOUS FORUM D. EUGENE STRANDNESS JR., M.D. MEMORIAL LECTURE

Each year, the American Venous Forum recognizes the significant contributions of an individual in research, education or clinical investigation in the field of venous diseases. The recipient of this distinction, chosen by the President of the American Venous Forum and confirmed by the Forum's Executive Committee, is named to the position of D. Eugene Strandness Jr., M.D. Memorial Lecturer and serves as the Keynote Speaker on a topic of his or her choice at the Annual Meeting of the Forum.

This honor, the highest given by the organization, has been bestowed to the following outstanding candidates in past years:

- 2007 Robert L. Kistner, M.D., Honolulu, Hawaii "Foresight 2020: Creating the Venous Vision" 2006 Pan Ganauly, Ph.D., Bethesda, MD "The Challenges in Venous Thrombosis" Michel R. Perrin, M.D., Chassieu, France 2005 "The Importance of International collaboration for the Development of a Scientific Approach to Venous Disease" 2004 Professor Eberhard Rabe, M.D., Bonn, Germany "Prevalence and Risk Factors of Chronic Venous Diseases: The Bonn Vein Study" 2003 Professor Claudio Allegra, M.D., Rome, Italy "Involvement of the Microcirculation in Chronic Venous Insufficiency" Professor Alfred Bollinger, M.D., Professor Emeritus, University 2002 of Zurich "Microcirculation in Chronic Venous Insufficiency and Lymphedema" 2001 Professor C.V. Ruckley, M.D., Edinburah, Scotland "Chronic Venous Insufficiency: Lessons from Scotland" Professor Sir Norman Browse, M.D., F.R.C.S., F.R.C.P. 2000 "Forty Years On" 1999 David Robinson, PhD, Bethesda, Maryland "A Journey to Complexity: The Continuing Evolution in Vascular Research"
- 1998 David Bergquist, M.D., Ph.D., Uppsala, Sweden "A Chronic Leg Ulcer - The Impact of Venous Disease"

- 1997 Professor Kevin G. Burnand, London, England "A Venous Thrombogene is and Thrombolysis"
- 1996 Ermenegildo A. Enrici, M.D., Buenos Aires, Argentina "The Role of the Perforants' System in Deep Venous Chronic Insufficiency in its Different Stages: Surgical Indications, Tactics and Techniques"
- 1995 Philip D. Coleridge Smith, M.D., FRCS, London, England "Venous Disease and Leukocyte Mediated Microcirculatory Injury"
- 1994 Andrew W. Nicolaides, M.D., FRCS, London, England "Deep Vein Thrombosis: Aetiology and Prevention. The Legacies of the 70's, The promises of the 80's and the Challenges of the 90's"
- 1993 Olav Thulesius, M.D., Ph.D., Linkoping, Sweden "Vein Wall Characteristics and Valvular Functions in Chronic Venous Insufficiency"
- 1992 G. W. Schmid-Schonbein, M.D., La Jolla, California "Leukocytes as Mediators of Tissue Injury"
- 1991 Jack Hirsh, M.D., Hamilton, Ontario, Canada "Development of Low Molecular Weight Heparin for Clinical Use"
- Hugo Partsch, M.D., Vienna, Austria"Diagnosis of AV Fistulas in Vascular Malformations"

AMERICAN VENOUS FORUM FOUNDATION RESEARCH AWARD

Each year The American Venous Forum Foundation offers a cash prize for up to three (3) abstracts on clinical or experimental work in venous diseases performed by residents in training, fellows and young physicians and surgeons in practice for less than five years



THE BSN-JOBST RESEARCH FELLOWSHIP IN VENOUS AND LYMPHATIC DISEASE

In 1995, The American Venous Forum Foundation announced the establishment of the BSN-Jobst, Inc. Research Fellowship in Venous and Lymphatic Disease.

- 1995 Peter J. Papas, M.D., UMDNJ New Jersey Medical School
- 1996 Jae-Sung Cho, M.D., Mayo Clinic, Rochester, MN
- 1997 Andrew C. Stanley, M.D., Burlington, VT
- 1998 Klaus See-Tho, M.D., Stanford University Medical Center
- 1999 Joseph D. Raffetto, M.D., Boston Medical Center
- 2000 No Award Given
- 2001 Brajesh K. Lal, M.D., UMDNJ New Jersey Medical School
- 2002 Susan O'Shea, M.D., Duke University Medical Center
- 2003 Charles Fields, M.D., Mayo Clinic
- 2004 John Rectenwald, M.D., University of Michigan
- 2005 Allesandra Puggioni, M.D., Mayo Clinic
- 2006 Stephanie K. Beidler, M.D., University of North Carolina
- 2007 Danny Vo, M.D., Mayo Clinic

The BSN-Jobst, Inc. Research Fellowship provides a one-year, \$25,000 grant to a research fellow chosen through a competitive peer-review selection process. A committee of distinguished vascular physicians, appointed by the American Venous Forum Foundation, determines the fellowship recipient and announces its selection during the Forum Finale.



SIGVARIS, INC. TRAVELING FELLOWSHIP IN VENOUS DISEASE

Sigvaris, Inc. initially established this \$12,000 Traveling Fellowship to provide a selected candidate with the opportunity to visit medical centers throughout the United States, Europe and elsewhere which have established themselves as centers of excellence in the management of venous disease. In 2006, the Award criteria was changed to encourage fellows to submit abstracts, and attend the Forum's Annual Meeting, and broadened to include up to four (4) finalists, who would each receive up to \$3,000 in travel reimbursement associated with attending the meeting. Finalists also receive free one-year candidate membership in the American Venous Forum.

1997	Mark H. Meissner, M.D., University of Washington Medical Center
1998	Paul R. Cordts, M.D., Triple Army Medical Center
1999	E. John Harris, Jr., M.D., Stanford University Medical Center
2000	Harold J. Welch, M.D., Lahey Clinic Medical Center
2001	David L. Gillespie, M.D., Uniformed Services University of the Health Sciences
2002	Joseph D. Raffetto, M.D., Boston Medical Center
2003	Audra Noel, M.D., Mayo Clinic
2004	Robert McLafferty, M.D., Southern Illinois University
2005	Antonios P. Gasparis, M.D., Stony Brook University
2006	Beverly Sharp, M.D., Charing Cross Hospital Biju Aravind, M.D., Charing Cross Hospital
2007	Alisha Oropallo, M.D., Boston Medical Center M. K. Barsoumi, M.D., Mayo Clinic Prandath Lall, M.D., Mayo Clinic Eugene Palchick, M.D., University of Rochester



The Servier Traveling Fellowship provides two fellows an opportunity to travel to France and Spain for the 2008 European Venous Forum to present his or her scientific research. Up to four (4) finalists are identified through a competitive peer-review process, and are invited to present their science during the AVF Meeting. Travel and accommodations for the finalists are reimbursed as part of the grant. The finalists are judged by an appointed AVF committee. Two winners will be selected to present their work at the European Venous Forum.

2006

Charles Stonerock, M.D., Indiana University School of Medicine Gustavo Oderich, M.D., Mayo Clinic

2007

Brian Knipp, M.D., University of Michigan Reagan Quan, M.D., Walter Reed Army Medical Center

BEST POSTERS

Each year, a formal poster session is held where authors are invited to give a 3-minute synopsis of their work followed by a 2-minute Q & A with the audience in attendance. Posters are scored and prizes are awarded to the top presentations.

2007 Winners:

Hemodynamic and Clinical Impact of the Lateral Embryonic Vein In Limps With Klippel-Trenaunay Syndrome

K. Delis, P. Gloviczki, P. Wennberg, T. Rooke, D.J. Driscoll – Mayo Clinic, Rochester, MN

Abdomino Pelvic Venous Assessment With Duplex Ultrasound (Transvaginal and Transparietal)

A. Sanchez, J. Leal, S. Zubicoa, L. Del Campo, F. Sainz – Hospital Ruber Internacional, Madrid, Spain

Impaired Cerebral Venous Hemodynamics In Multiple Sclerosis Patients

P. Zamboni, E. Menegatti, A. Legnaro, S. Gianesini, E. Fainardi, A. Liboni – University of Ferrara, Ferrara, Italy

GENERAL INFORMATION

REGISTRATION DESK

The Registration Desk will be located in the Gazebo (Charleston Place) and will be open during the following hours:

Tuesday, February 19 Wednesday, February 20 Thursday, February 21 Friday, February 22 Saturday, February 23 2:00 pm - 6:00 pm 7:00 am - 6:00 pm 7:00 am - 6:00 pm 7:00 am - 12:30 pm 7:00 am - 4:00 pm

REGISTRATION INFORMATION

Full Registration Fee Includes: All scientific sessions, Postgraduate Course, continental breakfast, coffee breaks and boxed lunches. In addition, the registration fee includes entrance to the Exhibit Hall, the Welcome Reception on Wednesday and the Forum Finale on Saturday evening.

Guest/Spouse Registration Fee Includes: The Welcome Reception, continental breakfast, mid-morning refreshments daily in the Hospitality Suite and the Forum Finale on Saturday evening.

ANNUAL BUSINESS MEETING LUNCH (Members Only)

The Annual Business Meeting will be held on Friday, February 22, 2008 at 12:00 pm in the Riviera Ballroom.

SPECIAL NEEDS

If you have a disability that requires special accommodations or assistance, please contact the AVF Administrative Offices prior



to the start of the Annual Meeting. Please advise the AVF Administrative Offices if you have any food allergies or dietary restrictions prior to the start of the Annual Meeting.

INSTRUCTIONS TO AUTHORS

Audio Visual. All presentations must be formatted using Power-Point. All presenters must bring their PowerPoint presentations on CD Rom or Flash Drive (USB) to the Speaker Ready Room at least 2-hours prior to their scheduled presentation.

Manuscripts. The American Venous Forum has a publication agreement covering papers from this meeting. Therefore, presenting authors of oral presentations **must** submit the full manuscript to the Journal of Vascular Surgery within 30-days of presentation. You must conform strictly with their guidelines when preparing your manuscript. All submissions to the Journal of Vascular Surgery must be made before the presentation date.

INDUSTRY PARTNERS

A select group of companies have been invited to participate in the Annual Meeting through sponsored symposia and exhibits. Sponsored symposia are listed in the program. The exhibits will feature the latest products for diagnosis and treatment of venous disease. Exhibits will be located in the Live Oak/Cypress/Dogwood Ballrooms. Continental breakfasts and refreshment breaks will be scheduled in the Exhibit Hall to give participants an opportunity to visit the various companies in attendance.

The Executive Committee and members of the American Venous Forum are most grateful to the following companies for their support of the Annual Meeting.

PLATINUM

Angiodynamics Jobst Juzo Sanofi-Aventis

GOLD

Bacchus Vascular Sigvaris, Inc. VNUS Medical Technologies

SILVER

Cook Medical Diomed, Inc.

BRONZE

Boston Scientific CircAid Medical Products Eisai, Inc. EKOS Corp. GE Healthcare Healthpoint, Ltd. Organogenesis, Inc. Possis Medical, Inc. Sonosite Terason Ultrasound



AMERICAN VENOUS FORUM

20th ANNUAL MEETING February 20-23, 2008

Charleston Place, Charleston, South Carolina

WEDNESDAY, FEBRUARY 20, 2008

7:00 am

Continental Breakfast

7:30 am POSTGRADUATE COURSE

IN THE ERA OF PAY FOR PERFORMANCE OUTCOMES ASSESSMENT IN VENOUS DISEASE: MEASUREMENT TOOLS AND RESULTS REPORTING

Educational Objectives:

At the conclusion of the Postgraduate Course, the attendees will be able to:

1. Identify outcome tools and measurements

- 2.Assess the results of treatment for venal caval filters, DVT, lymphedema and ulcer therapy
- 3.Assess compression devices and garments, clot extraction and superficial venous reflux

Additionally, databases and information systems, non-invasive testing and ICAVL reporting standards, severity of illness assessment tools and quality of life measurements will be presented. The participant will also gain an understanding of the international training lequirements and recommendations for the non-vascular specialist.

Session I

Introduction Joann Lohr, MD

Vena Caval Filters: Fact or Fiction David L. Gillespie, MD

Deep Venous Reconstruction: Stents and Valves *Peter Neglen, MD*

Lymphedema and Ulcer Therapy: Compressions Devices and Garments Joseph Raffetto, MD

Noninvasive Testing and ICAVL Reporting Standards Eugene R. Zierler, MD

Vascular Training Requirements and Recommendations For Nonvascular Specialists Bo Eklof, MD

Question and Answers & Discussion

9:30 - 10:00 am Coffee Break

Session II

Severity of Illness Assessment Tools and Quality of Life Michael Vasquez, MD

Thrombolysis – Who, What, When, Where, How For Clot Removal

Anthony Comerota, MD

Superficial Venous Reflux, Compression, and Nonoperative Therapy Marc A. Passman, MD

Superficial Venous Reflux Interventional Options Nick Morrison, MD

Databases and Information Systems Brenda K. Zierler, PhD

Question and Answer

12:00pm Conclusion

12:00 pm Lunch (Boxed Lunch To Be Provided)

1:00 pm		PROCEDU	
		Moderator:	Peter Neglen, MD
		Educational O	bjectives:
		1.To be aware	of technical aspects of venous stenting
		2.To realize it r	ole in chronic venous disease
		3.To recognize	e its adjuvant role in early clot removal
		Panelists:	Haraldur Bjarnason, MD David Gillespie, MD Anthony Gasparis, MD
3:00 pm		Coffee Bre	ak
3:30 pm			C SESSION I: Endovenous Mark Meissner, MD Lowell Kabnick, MD
		Educational O	bjectives:
		1.Upon comp	letion of this session attendees will understand:
			analysis of endovenous ablative therapies comparing pen procedures.
			uality of life and disease severity outcomes when superficial system with concomitant deep system reflux.
		,	f endovenous ablative procedures in patients that e had deep venous thrombosis.
			here is a benefit in performing immediate or staged phlebectomy with endovenous ablation.
			ice of inflammatory markers in venous ulcer, and the affect on therapy on changes in cytokines levels and healing.
			nce of foam sclerotherapy in treating unusual varices c nerve utilizing Duplex ultrasound guidance.
3:30 pm	1.	For Varicose S. Subramon Nottingham	ency Ablation Versus Conventional Surgery e Veins - A Comparison of Costs i ^{a1} , T. Lees ² - ¹ Queen's Medical Centre, , United Kingdom, ² Freeman Hospital, Newcastle Jnited Kingdom
3:50 pm	2.	Outcomes Venous Inst B. S. Knipp, F Blackburn, G	Is Laser Ablation Improves Venous Irrespective of the Presence of Deep ufficiency Mansoor, M. Hong, J. Bloom, E. Fellows, S. G. Adams, J. Pfeifer, D. Williams, T. Wakefield - Michigan, Ann Arbor, Ml
4:10 pm	3.	A. Puggioni,	Ablation of the Great Saphenous Vein In h Previous Venous Thrombosis: Is It Safe? N. Marks, A. Hingorani, A. Shiferson, E. Ascher as Medical Center, Brooklyn, NY

4:30 pm	4.	Mid-Term Results of the Surgical Treatment of Varices By Phlebectomy With Conservation of A Refluxing Saphenous Vein P. Pittaluga ¹ , S. Chastanet ¹ , J. J. Guex ² - ¹ Riviera Veine Institut,
		Nice, France; ² Cabinet de Medecine Vasculaire, Nice, France
4 :50 pm	5.	Endovenous Laser Therapy With Concomitant Or Sequential Phlebectomy: A Randomized Controlled Trial A. I. Mekako, J. Hatfield, M. N. Abdul Rahman, S. Gulati, P. T. McCollum, I. C. Chetter - Hull Royal Infirmary/University of Hull, Hull, United Kingdom
		MINI PRESENTATIONS
5:10 pm	6.	(Mini Presentation 1) Foam Sclerotherapy of Venous Malformations V. Cheng – San Diego Vein Institute, Encinitas, CA
5:15 pm	7.	(Mini Presentation 2) The Echo-Guided Sclerotherapy In Sciatic Nerve Varices Treatment S. Gianesini, G. Tacconi, A. Palazzo, P. Fortini, E. Righi, E. Menegatti, A. Liboni, P. Zamboni - Ferrara University, Ferrara, Italy
6:00 pm		Welcome Recepition

THURSDAY, FEBRUARY 21, 2008

7:00 am		Continental Breakfast / Exhibits Open
8:00 am		SCIENTIFIC SESSION II: Basic Moderators: Joseph Raffetto, MD & David L. Gillespie, MD
		Educational Objectives:
		1.Upon completion of this session attendees will gain knowledge in:
		2. The importance of microparticles in the formation of venous thrombosis.
		3.The influence of matrix metalloproteinases (MMPs) in the venous ulcer wound environment, and how the composition of MMPs may affect venous ulcer wound healing.
		 The importance of physiologic age on thrombus formation and resolution.
		5. How MMPs affect venous relaxation by altering the calcium entry into smooth muscle.
		Important novel hemodynamic measurements in defining chronic venous disease.
		7.Neovascularization and a novel treatment algorithm.
		8.Determining if cryo-venous stripping has any advantages over traditional stripping of the great saphenous vein.
8:00 am	8.	Microparticles Surface Proteins Influence Venous Thrombogenesis
		N. M. Abdullah, M. Kachman, A. Walker, A. E. Hawley, S. K. Wrobleski, D. D. Myers, Jr., J. R. Strahler, P. C. Andrews, P. K. Henke, T. W. Wakefield - University of Michigan, Ann Arbor, MI
8 :20 am	9.	Inflammatory Cytokine Levels In Chronic Venous Insufficiency Ulcers Before and After Compression Therapy
		S. Beidler, C. Douillet, D. Berndt, P. Rich, W. Marston - University of North Carolina at Chapel Hill, Chapel Hill, NC
8:40 am	10.	The Matrix Metalloproteinase (MMP) Profile In the Venous Ulcer Bed May Provide A Prognostic Indication of Ulcer Healing J. Tan, A. Smith, K. Burnand - Academic department of
		Surgery, Cardiovascular Division, London, United Kingdom
9:00 am	11.	The Prothrombotic Effects of Aging On Acute Venous Thrombosis In A Rodent Model A. P. McDonald, T. R. Meier, A. E. Hawley, J. N. Thibert, D. M. Farris, S. K. Wrobleski, P. K. Henke, T. W. Wakefield, D. D. Myers, Jr University of Michigan, Ann Arbor, MI

9:20 am	12.	MMP-2 Induced Venous Relaxation Via Inhibition of Ca2+ Entry-Dependent Mechanisms of Venous Smooth Muscle Contraction J. D. Raffetto ¹ , R. Khalil ² - ¹ VA Boston Healthcare System, West
		Roxbury, MA, ² Brigham and Women's Hospital, Boston, MA
		MINI PRESENTATIONS
9:40 am	13	(Mini Presentation 3)
		Doppler Derived Maximum Venous Outflow Velocity (MVOV) Demonstrates Asymmetric Lower Extremity Venous Flow In Normal Individuals M. Lebow, D. Cassada, O. Grandas, S. Stevens, M. Freeman, M. Goldman - UT Knoxville, Knoxville, TN
9:45 am	14	(Mini Presentation 4)
		Neovascularity and It's Treatment After Saphenous Ligation R. Bush, K. Hammond - Midwest Vein and Laser Center, Dayton, OH
9:50 am	15	(Mini Presentation 5)
		Cryo Strip Versus Classic Strip of the Great Saphenous Vein T. M. A. L. Klem ¹ , J. M. Schnater ² , P. R. Schutte ³ , A. C. van der Ham ⁴ , C. H. A. Wittens ⁵ - ¹ Erasmus Medical Center, Rotterdam, Netherlands; ² Academisch Medical Center, Amsterdam, Netherlands; ³ Albert Schweitzer Hospital, Dordrecht, Netherlands; ⁴ Sint Franciscus Hospital, Rotterdam, Netherlands; ⁵ Haga Hospital, The Hague, Netherlands
10:00 am		Coffee Break
10:30 am		SCIENTIFIC SESSION III: FOAM & DIAGNOSTICS Moderator: Peter Pappas, MD
		Educational Objectives:
		1. Upon completion of this session attendees will understand:
		If positioning of the patient undergoing foam sclerotherapy is important in preventing foam particle central migration.
		3. The application of Duplex ultrasound and MRI in understanding the mechanism of pneumatic compression on the venous and muscles of the lower extremity.
		4.The advantages or disadvantages in a randomized trial comparing treatment of varicose veins by surgery versus foam sclerotherapy.
		5. The objectives of the National Venous Screening Program, how the NVSP has impacted the perception and education of the public and practitioners, and future direction in establishing nation-wide screening with emphasis on, risk of venous thromboembolism, venous clinical severity, primary care education, and timely specialty care referral and treatment.

		6.The changes of MMPs and its naturally occurring inhibitors in the venous wall of varicose veins, and how this may impact on pathophysiology.
		 Demographic and risk factors that can effect pain following endovenous ablation.
10:30 am	16	Assessment of Techniques To Reduce Sclerosant Foam Migration During Ultrasound Guided Sclerotherapy D. A. Hill, R. Hamilton – The Vein Treatment Centre, Calgary, AB, Canada
10:50 am	17	Combined MRI and Duplex Ultrasound Investigation of the Mechanism of Action of the Pneumatic Compression Devices F. Lurie ¹ , H. Yoon ² , V. Scott ³ , R. L. Kistner ¹ – ¹ University of Hawaii and Kistner Vein Clinic, Honolulu, HI, ² Hawaii Permanente Medical Group, Inc., Honolulu, HI, ³ Keck School of Medicine USC, Los Angeles, CA
11:10 am	18	Comparison Between Surgical Treatment and Ultrasound-Guided Microfoam Sclerotherapy For Patients With Primary Varicose Veins In the Lower Limbs: Early Results of A Randomized Controlled Trial M. Figueiredo, S. P. Araujo, F. Miranda Jr Escola Paulista de Medicina - Unifesp, Uberlandia, Brazil
		MINI PRESENTATIONS
11:30 am	19	(Mini Presentation 6)
		National Venous Screening Program – An Update Marc Passman, MD
	20	WITHDRAWN
11:35 am	21	(Mini Presentation 8)
		Patient Characteristics and Treatment Factors That Affect Pain Following Endovenous Laser Treatment (EVLT) For Venous Insufficiency P. A. Hertzman, B. Peterson ² - ¹ Vein Care of New Mexico, Los Alamos, NM, ² University of New Mexico, Albuqueque, NM

12:00 pm	AMERICAN COLLEGE OF PHLEBOLOGY SCLEROTHERAPY SESSION Moderator: Steve Zimmet, MD & Nick Morrison, MD
	Educational Objectives:
	1.Better utilize sclerotherapy to treat incompetent varices
	2.Minimize risk of complications of sclerotherapyy
	3.Understand issues related to the importation, compounding and off-label use of sclerosants
12:00 pm	Office Set-Up and Sclerotherapy Techniques Nick Morrison, MD
12:15 pm	Sclerotherapy: Cleaning Up Before & After Endovenous Laser Robert Min, MD
12:30 pm	X-Ray Guided Sclerotherapy Mel Rosenblatt, MD
12:45 pm	Sclerosants: Importation, Compouding and Off- Label Use Steve Zimmet, MD
1:30-5:50 pm	INDUSTRY WORKSHOPS (Three 80-Minute Sessions)
	Ultrasound Investigations for Venous Disease Moderator: Nicos Labropoulos, MD
	Educational Objectives:
	 Understanding of basic normal venous anatomy identified by venous ultrasound.
	2. Understanding of diagnostic criteria for venous thrombosis using venous ultrasound.
	3. Understanding of diagnostic criteria for venous insufficiency (deep, superficial, perforator) using venous ultrasound
	Endovenous Ablation of the Saphenous Vein Moderator: Michael Vasquez, MD
	Educational Objectives:
	 Understand and perform U/S guided access of an enlarged saphenous vein based on practice on a model.
	2.Discuss the importance and technique of intra-compartmental tumescent anesthesia for the performance of endovenous saphenous vein ablation.
	3. Identify different modalities of saphenous vein ablation for possible integration into their practice.

Pharmaco-Mechanical Thrombectomy (PMT)

Moderator: Peter Lin, MD

Educational Objectives:

- 1. Understand endovascular treatment strategies of acute deep venous thrombosis.
- 2. Understand the role of mechanical thrombectomy in the treatment of acute deep venous thrombosis
- 3. Be familiar with various mechanical thrombectomy devices in the treatment of acute deep venous thrombosis.
- 4. Have insight into potential applications of pharmacomechanical thrombectomy in acute deep venous thrombosis.

Venous Ulcer Wound Care

Moderator: William Marston, MD

Educational Objectives:

- 1. Evaluate various methods of compression and the advantages and disadvantages of each in the treatment of venous leg ulcers
- 2. Consider the vast range of products available to apply to the wound surface of venous leg ulcers and learn strategies to choose the best ones for each leg ulcer
- 3. Review the active therapies available that are proven to accelerate the healing of leg ulcers and demonstrate proper application techniques for these products

OR CONCURRENT SYMPOSIUM SESSION

1:30 – 2:50 pm	Venous Coding and Maximizing Reimbursement Moderator: Robert Zwolak, MD
	Educational Objectives:
	 Use appropriate category one CPT codes to report standard venous operations
	2.Understand the requirements for development of new CPT codes
	 Have a working familiarity with the method by which CPT codes are valued
3:00 – 4:20 pm	Venographic Assessment Moderator: David Gillespie, MD
	Educational Objectives:
	 Understand the indications and techniques for performing ascending venography
	2.Understand the indications and techniques for performing ilio/ cavography
	 Understand the indications and techniques for performing ovarian vein/pelvic dumping imaging
3:00 – 3:15 pm	Ascending/Decending Venography David Gillespie, MD

3:15 – 3:30 pm	Venographic Assessment of PelvicCongestion Syndrome Mark Meissner, MD
3:30 – 3:45 pm	Extremity Venography For Venous TOS Marc Passman, MD
4:30 – 5:50 pm	What To Do With Recurrent Varicose Veins? Moderator: Andre van Rij, MD
	Educational Objectives: 1.Understand the causes of recurrence of varicose veins and the role that neovascularisation has in this
	 Gain a basic understanding of the biology of neovascularisation, and recanalisation,
	 Be aware of how this varies with different treatments of varicose veins and how it might be prevented.
	4.Be familiar with treatments for recurrence and their relative merit.
	5. Have a rationale for counseling patients regarding the risk of recurrence following varicose vein treatment.
	PLEASE NOTE: The following evening symposium is included in the registration fee for physicians and allied health professionals. However, seating is limited and pre-registration is required. We regret that due to strict codes, spouses and guests may not attend.
6:30 – 8:30 pm	EVENING SYMPOSIUM Supported by Bacchus Vascular and Sanofi Aventis.
6:30 – 8:30 pm	
6:30 – 8:30 pm	Supported by Bacchus Vascular and Sanofi Aventis. THE TIMES THEY ARE A-CHANGING: VENOUS
6:30 – 8:30 pm	Supported by Bacchus Vascular and Sanofi Aventis. THE TIMES THEY ARE A-CHANGING: VENOUS THROMBOEMBOLISM UPDATE 2008 Educational Objectives:
6:30 – 8:30 pm	Supported by Bacchus Vascular and Sanofi Aventis. THE TIMES THEY ARE A-CHANGING: VENOUS THROMBOEMBOLISM UPDATE 2008 Educational Objectives: 1.Be familiar with the latest ACCP Chest Guidelines 2.Understand current concepts regarding the treatment choices
6:30 – 8:30 pm 6:30 – 6:50 pm	Supported by Bacchus Vascular and Sanofi Aventis. THE TIMES THEY ARE A-CHANGING: VENOUS THROMBOEMBOLISM UPDATE 2008 Educational Objectives: 1.Be familiar with the latest ACCP Chest Guidelines 2.Understand current concepts regarding the treatment choices and duration of treatment for venous thromboembolism 3.Understand the differences between iliofemoral thrombosis and
	Supported by Bacchus Vascular and Sanofi Aventis. THE TIMES THEY ARE A-CHANGING: VENOUS THROMBOEMBOLISM UPDATE 2008 Educational Objectives: 1.Be familiar with the latest ACCP Chest Guidelines 2.Understand current concepts regarding the treatment choices and duration of treatment for venous thromboembolism 3.Understand the differences between illofemoral thrombosis and other forms of venous thrombosis Putting New Joint Commission Quality Standards For DVT Into Hospital Practice

7:30 – 7:50 pm	Iliofemoral DVT Thrombus Removal Techniques: Safe and Effective Michael Zatina, MD
7:50 – 8:10 pm	Mechanolytic Intervention For Iliofemoral DVT and the Need For A RCT Anthony Comerota, MD
8:10 – 8:30 pm	Panel Discussion/Q&A

FRIDAY, FEBRUARY 22, 2008

7:00 am		Continental Breakfast / Exhibits Open
7:30 am		SCIENTIFIC SESSION IV: CHRONIC VENOUS DISEASE Moderators: Michael Ricci, MD & Marc Passman, MD
		Educational Objectives:
		1.Upon completion of this session attendees will be able to:
		 Understand the variability in venous Duplex laboratory reporting and the need for uniformity.
		3.Understand the important application of intravascular ultrasound in placement of vena cava filters in multi-trauma patients with emphasis on technique, anatomic variations, complications, and early durability.
		4.A novel technique utilizing an endovenous valve stent for treating patients with deep venous insufficiency.
		 Understand the hemodynamic relation between the iliac venous system and the saphenofemoral junction.
		6.Appreciate the complex nature of varicose vein formation and chronic venous disease, and the role that hypoxia may have on its pathogenesis.
7:30 am	22	Use of A Structured Audit Tool In Assessing Venous Duplex Reports D. L. Wooster - University of Toronto, Toronto, ON, Canada
7:50 am	23	Intravascular Ultrasound Guided Inferior Vena Cava Filter Placement In the Multi-Trauma Patients From Global War On Terrorism: A Single Center Experience G. Aidinian, A. A. Amin, P. W. White, E. Adams, C. J. Fox, M. Cox, D. L. Gillespie - Walter Reed Army Medical Center, Washington, DC
8:10 am	24	The Effect of Pressure On Migration and Further Characterization of the Venous Ulcer Fibroblast G. Scriver, A. Stanley, M. A. Ricci, K. Corrow, M. Slusarczyk, S. Shackford, J. Adams, G. Steinthorsson, D. Berges, A. Howard - University of Vermont, Burlington, VT
8:30 am		EUROPEAN VENOUS FORUM – FIRST PLACE WINNER
		Haemodynamic Assessment of Iliac Veins and Their Relation With the Sapheno-Femoral Junction P. Brazis ¹ , R. Piotrowicz ¹ , N. Labropoulos ² , A. Jawien ¹ ¹ Department of Surgery, Collegium Medicum, Nicolaus Copernicus University, Bydgoszcz, Poland - ² Division of Vascular Surgery, University of Medicine & Dentistry of New Jersey New Jersey Medical School, Newark, NJ USA

8:40 am		EUROPEAN VENOUS FORUM – SECOND PLACE WINNER
		Is Hypoxia A Feature of Varicose Vein Disease?
		B. Sharp ^{1,2} , B. T. Navin ¹ , C. Monaco ^{1,2} , E. Paleolog ^{1,2} , A. H. Davies ² - ¹ Kennedy Institute of Rheumatology, Imperial College, London, UK; ² Charing Cross Hospital Department of Surgery, Oncology and Anaesthetics (SORA), Imperial College, London, UK
9:10 am		Coffee Break / Visit Exhibits
9:40 am		SCIENTIFIC SESSION V: ENDOVENOUS STENTING Moderators: Robert McLafferty, MD & Peter Neglen, MD
		Educational Objectives:
		1.Upon completion of this session attendees will be able to:
		2. Understand the variability in venous Duplex laboratory reporting and the need for uniformity.
		3. Understand the important application of intravascular ultrasound in placement of vena cava filters in multi-trauma patients with emphasis on technique, anatomic variations, complications, and early durability.
		4.A novel technique utilizing an endovenous valve stent for treating patients with deep venous insufficiency.
		5. Understand the hemodynamic relation between the iliac venous system and the saphenofemoral junction.
		6. Appreciate the complex nature of varicose vein formation and chronic venous disease, and the role that hypoxia may have on its pathogenesis.
9:40 am	25	Venous Stenting Across the Inguinal Ligament P. Neglén, P. Tackett, S. Raju - River Oaks Hospital, Flowood, MS
10:00 am	26	Reinterventions After Venous Stenting For Chronic Venous Disease S. Raju ¹ , P. Tackett ² , P. Neglén ² - ¹ University of Mississippi Medical Center, Jackson, MS, ² River Oaks Hospital, Flowood, MS
10:20 am	27	WITHDRAWN
10:20 am		Venous Stent Registry Update BK Lal, MD
10:25 am		AVF Update – Where the Forum Is Going Mark H. Meissner, MD
10:30 am		Founders Award (TBA) Presented By: Mark H. Meissner, MD

10:35 am	2007 Award Update Introduced By: Mark H. Meissner, MD
	2007 BSN Jobst Winner – Report Danny Vo, MD, Mayo Clinic
10:45 am	2007 Servier Traveling Fellowship Winners – Report Brian Knipp, MD, University of Michigan
10:55 am	2007 Sigvaris Fellowship – Announcement David Gillespie, MD for Reagan Quan, MD, Walter Reed Army Medical Center
11:00 am	PRESIDENTIAL ADDRESS Mark H. Meissner, MD Introduction By: Joann M. Lohr, MD
12:00 pm	MEMBER BUSINESS LUNCH
Free Afternoon	Golf & Tennis

SATURDAY, FEBRUARY 23, 2008

7:00 am		Continental Breakfast – Visit Exhibits
8:00 am		SCIENTIFIC SESSION VI: VENOUS THROMBOEMBOLISM Moderators: Joseph Caprini, MD & Peter Henke, MD
		Educational Objectives:
		1.Upon completion of this session attendees will be able to:
		 Understand the utilization of Duplex ultrasound in defining temporal changes of venous thrombi.
		3.Understand the potential usefulness of combining d-dimer and lower extremity Duplex ultrasound tests in predicting outcome for venous thromboembolism in high risk patients undergoing surgery
		 Understand the complex and controversial management of pregnant females that had previous ilio-caval stenting.
		5.Understand the benefits and limitations of diagnostic modalities in patients with suspected venous thromboembolism
		6.Understand the indications, strategy, technical aspects, pharmacologic drugs, mechanical application, and outcomes in treating patients with iliofemoral deep venous thrombosis with pharmacomechanical thrombolysis.
8:00 am	28	Time-Course Analysis of Venous Thrombus With Ultrasonographic Tissue Elasticity Imaging - Preliminary Findings K. Uno ^{1,5} , A. Tonomura ² , T. Osaka ² , T. Mitake ² , M. Suda ³ , M. Yamakawa ³ , Y. Isaka ⁴ , S. Homma ⁵ , T. Shiina ³ , K. Aonuma ⁵ - ¹ Namegata District General Hospital, Namegata, Japan, ² Ultrasound Systems Division, Hitachi Medical Corporation, Kashiwa, Japan, ³ Graduate School of System and Information Engineering, University of Tsukuba, Tsukuba, Japan, ⁴ Medical Branch, Academic Service Office for Comprehensive Human Sciences, University of Tsukuba, Tsukuba, Japan, ⁵ Cardiovascular Division, Institute of Clinical Medicine, Graduate School of Comprehensive Human Science, University of Tsukuba, Tsukuba, Japan
8:20 am	29	Do Preoperative D-Dimer Testing and Venous Duplex Scanning of the Lower Extremities Alter the Outcome In Patients At High Risk For Postoperative Venous Thromboembolism? T. Yamaki ¹ , M. Nozaki ¹ , H. Sakurai ¹ , M. Takeuchi ² , K. Soejima ³ , T. Kono ¹ - ¹ Tokyo Women's Medical University, Tokyo, Japan, ² Nihon University, Tokyo, Japan, ³ Tokyo Metropolitan Hiroo General Hospital, Tokyo, Japan
8:40 am	30	Management of Pregnancy In Women With Previous Ilio-Caval Stenting O. Hartung - CHU Nord, Marseille, France

9:00 am	31	The Evaluation of Diagnostic Procedures of Venous Thromboembolism (VTE) In Patients With Suspected VTE J. Lee, B. Zierler - University of Washington, Seattle, WA
9:20 am	32	The Quantitative Benefit of Isolated, Segmental, Pharmacomechanical Thrombolysis For Iliofemoral DVT J. Martinez, A. J. Comerota, S. Kazanjian, R. DiSalle, D. M. Sepanski, Z. I. Assi - The Toledo Hospital, Toledo, OH
9:40 am		Coffee Break / Visit Exhibits
10:10 am		SCIENTIFIC SESSION VII: CHRONIC VENOUS DISEASE II Moderators: Joann Lohr, MD & Fedor Lurie, MD
		Educational Objectives:
		1.Upon completion of this session attendees will understand:
		 The implications of chronic venous disease and venous hypertension on the adverse effect on arterial hemodynamics.
		3. The risk factors implicated in patients with varicose veins that will progress to venous ulceration.
		4. The risk factors associated with recalcitrant venous ulcer treated with compression.
		5. The technical application of a neovalve for deep venous insufficiency, comparing two different experiences.
10:10 am	33	Lower Extremity Arterial Inflow Is Adversely Affected In Patients With Postthrombotic Venous Disease D. Paolini, L. Jones, A. J. Comerota - The Toledo Hospital, Toledo, OH
10:30 am	34	Which Patients With Varicose Veins Are At Increased Risk of Chronic Leg Ulceration? L. Robertson ¹ , A. Lee ² , K. Gallagher ³ , S. Carmichael ⁶ , C. Evans ⁴ , B. McKinstry ¹ , S. Fraser ³ , P. Allan ¹ , C. Ruckley ¹ , F. Fowkes ¹ - ¹ University of Edinburgh, Edinburgh, United Kingdom, ² University of Aberdeen, Aberdeen, United Kingdom, ³ Lothian University Hospitals NHS Trust, Edinburgh, United Kingdom, ⁴ NHS Lothian, Edinburgh, United Kingdom
10:50 am	35	Risk Factors Related To the Failure of Venous Leg Ulcers To Heal With Compression Treatment D. J. Milic, S. S. Zivic, D. C. Bogdanovic, V. D. Milojkovic, M.A. Pejic, V. M. Popovic - Clinic for Vascular Surgery, Clinical Centre Nis, Nis, Serbia
11:00 am	36	Neovalve Construction In Deep Venous Incompetence: Comparison Between Two Subsequent Case Series and Related Technical Details M. Lugli, S. Guerzoni, O. Maleti - Hesperia Hospital, Modena, Italy
11:30 am		D. EUGENE STRANDNESS MEMORIAL LECTURE Introduced by: Mark H. Meissner, MD

12:30 – 1:30 pm	INDUSTRY SPONSORED LUNCHEON
	ClosureFAST™ Clinical Trials Update
	By: VNUS Medical Technologies
	1) Prospective, multicenter 1 year follow-up
	2) Randomized trial comparing ClosureFAST to endovenous laser
	3) Lessons learned 1 year post launch
1:30 pm	ASK THE EXPERTS: PELVIC CONGESTION SYNDROME Moderator: Chieh Min Fan, MD
	Educational Objectives:
	 Review anatomy and techniques for imaging the venous structures of the pelvis and retroperitoneum.
	2.Recognize the clinical manifestations of pelvic venous congestion
	 Understand endovascular and surgical treatment approaches for pelvic venous congestion syndromes
	 Review classification, clinical patterns, and treatment approach for pelvic vascular anomalies
	Anatomy and Imaging of the Venous System of the Pelvis and Retroperitoneum Chieh Min Fan, MD, Brigham and Women's Hospital, Boston, MA
	Pelvic Congestion Syndrome: Diagnosis and
	Management Anthony C. Venbrux, MD, George Washington U. Hospital, Washington, DC
	Nutcracker Syndromes: Endovascular and Surgical Management Matthew Menard, MD, Brigham and Women's Hospital, Boston, MA
	Vascular Anomalies: An Uncommon Cause of Pelvic Venous Congestion Patricia Burrows, MD, St. Lukes – Roosevelt Hospital Center, New York, NY

2:30 pm Coffee Break / Visit Exhibits

HOW TO SESSION

Recanalization and Re-Endovenous Ablation; Mapping Out My Veins Tips and Tricks; Sclero the Do's and Don'ts

Moderator: Julianne Stoughton, MD

Educational Objectives:

- 1.Attendees will become familiar with many of the common, and some of the unusual (but important) patterns of venous anatomy
- 2.There will be a discussion involving the approach to incompetent perforating veins: reviewing the treatment options, the technical aspects of each treatment, as well as a discussion of which veins are best treated with which technology
- 3.Recannalization, neovascularization and recurrent veins after venous intervention and will be discussed. The approaches will be illustrated with case presentations
- 4.Difficult management cases will be presented including: the hypercoaguable patient, the obese patient, patients with anomalous anatomy, etc.

Endovenous Heat Induced Thrombosis: When To, How To and What To Look For

Lowell Kabnick, MD

Treatment of Incompetent Perforators Steve Elias, MD

Treatment of Neovascularization and Recannalized Veins

Ronald Bush, MD

4:00 pm	MODERATED POSTER SESSION Moderator: Michael Dalsing, MD Frank Padberg, MD Bo Eklof, MD
	Educational Objectives: The participants in the poster session will gain a wide range of knowledge expansion including chronic venous disorder, saphenous vein treatment, understanding risk factors and evaluation methods.
P-1	Microparticles: A Natural History Time Course Analysis In A Model of Murine Venous Thrombosis A. E. Hawley, D. M. Farris, N. E. Ballard, A. P. McDonald, S. K. Wrobleski, P. K. Henke, D. D. Myers, T. W. Wakefield - University of Michigan, Ann Arbor, Ml
P-2	Popliteal Vein Compression Syndrome: Obesity, Venous Disease and the Popliteal Connection R. J. Lane ¹ , M. L. Cuzilla ² - ¹ Royal North Shore Hospital, Sydney, Australia ² Vascular Surgery Inventigations and Managment, Sydney, Australia
P-3	Ultrasonic Venous Valve Imaging - A Prerequisite For Exostent Repair R. J. Lane ¹ , M. N. Phillips ² , M. L. Cuzilla ³ - ¹ Royal North Shore Hospital, Sydney, Australia; ² AllVascular Pty Ltd, Sydney, Australia; ³ Vascular Surgery Inventigations and Managment, Sydney, Australia
P-4	Prevalence and Distribution of Deep Vein Thrombosis In Patients With Symptomatic Pulmonary Embolism T. Yamaki ¹ , M. Nozaki ¹ , H. Sakurai ¹ , M. Takeuchi ² , K. Soejima ³ , T. Kono ¹ - ¹ Tokyo Women's Medical University, Tokyo, Japan; ² Nihon University, Tokyo, Japan; ³ Tokyo Metropolitan Hiroo General Hospotal, Tokyo, Japan
P-5	Endovenous Laser Therapy In the Treatment of Short Saphenous Varicose Veins: A Non-Randomised Controlled Trial A. Mekako, J. Hatfield, S. Gulati, M. Abdul Rahman, P. T. McCollum, I. C. Chetter - Hull Royal Infirmary/University of Hull, Hull, United Kingdom
P-6	Greater Saphenous Vein Diameter Predicts Venous Reflux J. Bloom, F. C. Vandy, S. Brown, A. Clay, C. Lane, G. Reynolds, S. LeBaron, C. Nighswander, P. K. Henke, T. W. Wakefield - University of Michigan, Ann Arbor, MI

P-7	Combined Intermittent Pneumatic Leg Compression and Pharmacological Prophylaxis For Prevention of Venous Thromboembolism In High Risk Patients S. K. Kakkos ¹ , J. A. Caprin ^P , G. Geroulakos ³ , A. N. Nicolaides ³ , G. P. Stansby ⁴ , D. J. Reddy ¹ - ¹ Henry Ford Hospital, Detroit, MI; ² Evanston Northwestern Healthcare, Evanston, IL; ³ Imperial College, London, United Kingdom; ⁴ University of Newcastle upon Tyne, Newcastle upon Tyne, United Kingdom
P-8	Evaluation of Venous Thromboembolism Prophylaxis In Randomly Selected Medical and Surgical Patients <i>B. K. Zierler, J. Lee, G. Han, H. Oh, C. Jacobson - University of</i> <i>Washington, Seattle, WA</i>
P-9	An Algorithm For Outpatient Deep Venous Thrombosis Management and Severe Post-Thrombotic Syndrome At Mid-Term Follow-Up E. Sivrikoz, M. Kurtoglu - Dept. of General Surgery, Istanbul School of Medicine, Istanbul University, Istanbul, Turkey
P-10	Prevalence of Isolated (C2) and Complicated (C2+) Varicose Veins Among Patients Consulting Vascular Specialists For Varicosis: A Snapshot M. Cazaubon ¹ , M. Lugli ² , P. Burseta ³ , M. Perrin ⁴ , F. A. Allaert, V ⁵ - ¹ American hospital, Neuilly, France; ² Vascular hospital, Bologna, Italy; ³ vascular dpt hopital, Modena, Italy; ⁴ Vascular dpt hospital, Lyon, France, ⁵ cenbiotech ceren ESC and DIM CHRU Dijon, Dijon, France
P-11	Prospective Randomized Efficacy of Ultrasound- Guided Foam Sclerotherapy Compared To Ultrasound-Guided Liquid Sclerotherapy In the Treatment of Symptomatic Venous Malformations T. Yamaki ¹ , M. Nozaki ¹ , H. Sakurai ¹ , M. Takeuchi ² , K. Soejima ³ , T. Kono ¹ - 1Tokyo Women's Medical University, Tokyo, Japan; ² Nihon University, Tokyo, Japan; ³ Tokyo Metropolitan Hiroo General Hospital, Tokyo, Japan
P-12	Elastic Stockings and Ulcer Treatment: What About Pressure and Stiffness? G. Mosti - Clinica MD Barbantini, Lucca (LU), Italy
P-13	Inelastic Compression Increases Venous Ejection Fraction More Than Elastic Bandages G. Mosti ¹ , H. Partsch ² , V. Mattaliano ¹ - 1Clinica MD Barbantini, Lucca (LU), Italy; ² Dermatology Department, Vienna, Austria
P-14	Arterial Revascularization and Compression Therapy In the Treatment of Mixed Arterial/Venous Leg Ulcers D. J. Milic, S. S. Zivic, Z. D. Perisic, R. Jankovic, G. Djordjevic, D.M. Stamenkovic, Z.D. Maksimovic Clinic for Vascular Surgery, Clinical Centre Nis, Serbia

P-15	Morphological Changes On Varicose Vein Wall Corresponds To MMP/TIMP Alterations B. Aravind ¹ , T. Navin ² , C. Monaco ² , E. Paleolog ² , A. H. Davies ¹ - ¹ Imperial College, Charing Cross Hospital, London, United Kingdom, ² Kennedy Institute of Rheumatology, Imperial College, London, United Kingdom
P-16	Lower Power Improves Clinical Outcome of the Endovenous Laser Treatment S. Kaspar, J. Siller, Z. Cervinkova - Flebocentrum, Hradec Kralove, Czech Republic
P-17	Effectiveness of Weight Loss On the Evolution of Chronic Venous Insufficiency (CVI) After Bariatric Surgery In Obese Patients J. Benigni ¹ , J. Uh ^p , J. Gobin ³ , A. Capella ⁴ - ¹ Hôpital BEGIN, St Mandé, France; ² Surgical Venous Center, Neuilly, France; ³ Vascular medecine, Lyon, France; ⁴ Angiologist, Paris, France
P-18	One Year Follow-Up of Radiofrequency Segmental Thermal Ablation (RTFA) of Great Saphenous Veins <i>T. M. Proebstle</i> ¹ , <i>B. Vago</i> ² , <i>J. ALm</i> ³ , <i>O. Goeckeritz4</i> , <i>C. Lebard</i> ⁵ , <i>O. Pichot6</i> - 1 <i>University of Mainz, Mainz, Germany;</i> ² <i>University</i> <i>of Heidelberg, Heidelberg, Germany;</i> ³ <i>Dermatoligicum,</i> <i>Hamburg, Germany;</i> ⁴ <i>Venenzentrum am Elsterpark, Leipzig,</i> <i>Germany;</i> ⁵ <i>Hospital St. Michel, Paris, France;</i> ⁶ <i>CHU Service de</i> <i>Chirurgie Vasculaire, Grenoble, France</i>
P-19	Elimination of Superficial Reflux With Or Without Subcutaneous Fasciotomy - The Impact On Deep Axial Reflux and ulcer Healing J. T. Christenson - Division of Cardiovascular Surgery, University Hospital of Geneva, Geneva, Switzerland
P-20	The Effectiveness and Use of Compression Stockings of Various Strength For the Treatment of Venous Disorders and Diseases: A Literature Survey W. Blaettler ¹ , H. E. Gerlach ² , F. Amsler ³ - ¹ Angio Bellaria, Zürich, Switzerland; ² Center for Vascular Diseases, Mannheim, Germany; ³ Amsler Consulting, Biel-Benken, Switzerland
P-21	Management of Venous Injuries At the Air Force Theater Hospital In Balad, Iraq S. Gifford ¹ , W. T. Jones ¹ , M. A. Ricc ² , W. D. Clouse ¹ , T. E. Rasmussen ¹ - ¹ Wilford Hall Medical Center, San Antonio, TX; ² University of Vermont, Burlington, VT
P-22	Critical Issues In the Management of Venous Malformation (VM) Coexisting With Lymphatic Malformation (LM) - Klippel Trenaunay Syndrome (KTS) B. Lee, J. Laredo, D. Deaton, R. Neville - Georgetown University, Washington, DC

P-23	Variability of Interface Pressure Exerted By Compression Bandages and Standard Size Compression Stockings H. Partsch ¹ , W. Vanscheidt ² - ¹ Medical University Vienna, Vienna, Austria; ² University Clinic for Dermatology, Freiburg i.Br., Germany
P-24	Recanalization of Short Saphenous Vein After EVLT S. Shokoku - Varix Ambulatory Surgery Center, Okayama Daiichi Hospital, Okayama-shi, Japan
P-25	Endovenous Laser Ablation Compared With Stripping - Multi-Center RCT In Japan T. Ogawa ¹ , S. Hoshino ¹ , S. Makimura ² , H. Shigematsu ² , N. Azuma ³ , T. Sasajima ³ , H. Sugawara ⁴ , M. Ichikt ⁴ , S. Shokoku ⁵ - ¹ Fukushima Dalichi Hospital, Fukushima, Japan; ² Tokyo Medical University, Tokyo, Japan; ³ Asahikawa Meidical College, Asahikawa, Japan; ⁴ JR Sendai Hospital, Sendai, Japan; ⁵ Okayama Dalichi Hospital, Okayama, Japan
P-26	Pulse*Spray Sclerotherapy Study: A Pilot Study J. I. Almeida, J. K. Raines - Miami Vein Center, Miami, FL
P-27	Incompetent Perforators - What We Think We Know P. A. Hertzman - Vein Care of New Mexico, Los Alamos, NM
P-28	Non-Saphenous Approach To Varicose Veins With Foam Sclerotherapy V. Cheng - San Diego Vein Institute, Encinitas, CA
P-29	Case Report: Epitheloid Hemangioendothelioma of the Common Femoral Vein M. Lebow, A. Hurd, D. Cassada, M. Freeman, O. Grandas, S. Stevens, M. Goldman - University of Tennessee, Knoxville, TN
P-30	A Report of Two Rare Cases of Venous Aneurysms Involving the Lesser Saphenous Venous System S. Chen, A. N. Bowser, W. D. Clouse, C. Johnson, T. E. Rasmussen - Wilford Hall Medical Center Lackland AFB, San Antonio, TX
P-31	Multimodal Endovascular - Open Surgical Approach To Phlegmasia Cerulea Dolens of the Upper Extremity: A Case Report N. Patel, A. Puggioni, X. Li, A. Hingorani, A. Shiferson, V. Tran, E. Ascher - Maimonides Medical Center, Brooklyn, NY
P-32	Relation Between Number of Pregnancies and Great Saphenous Vein Diameters S. X. Salles-Cunha ¹ , N. Morrison ² - ¹ CompuDiagnostics, Inc, Phoenix, AZ; ² Morrison Vein Institute, Phoenix, AZ
7:30 pm	THE FORUM FINALE Awards, Dinner, Entertainment & More

WEDNESDAY, FEBRUARY 20, 2008

7:00 am

ednesdav

7:30 am

Continental Breakfast

POSTGRADUATE COURSE

IN THE ERA OF PAY FOR PERFORMANCE OUTCOMES ASSESSMENT IN VENOUS DISEASE: MEASUREMENT TOOLS AND RESULTS REPORTING

Educational Objectives:

At the conclusion of the Postgraduate Course, the attendees will be able to:

1. Identify outcome tools and measurements

- 2.Assess the results of treatment for venal caval filters, DVT, lymphedema and ulcer therapy
- 3.Assess compression devices and garments, clot extraction and superficial venous reflux

Additionally, databases and information systems, non-invasive testing and ICAVL reporting standards, severity of illness assessment tools and quality of life measurements will be presented. The participant will also gain an understanding of the international training lequirements and recommendations for the non-vascular specialist.

Session I

Introduction Joann Lohr, MD

Vena Caval Filters: Fact or Fiction David L. Gillespie, MD

Deep Venous Reconstruction: Stents and Valves Peter Neglen, MD

Lymphedema and Ulcer Therapy: Compressions Devices and Garments Joseph Raffetto, MD

Noninvasive Testing and ICAVL Reporting Standards

Eugene R. Zierler, MD

Vascular Training Requirements and Recommendations For Nonvascular Specialists Bo Eklof, MD

Question and Answers & Discussion

9:30 - 10:00 am Coffee Break

Session II

	Severity of Illness Assessment Tools and Quality of Life Michael Vasquez, MD	e
	Thrombolysis – Who, What, When, Where, How For Cle Removal Anthony Comerota, MD	ot
	Superficial Venous Reflux, Compression, and Nonoperative Therapy Marc A. Passman, MD	
	Superficial Venous Reflux Interventional Options Nick Morrison, MD	
	Databases and Information Systems Brenda K. Zierler, PhD	
	Question and Answer	
12:00pm	Conclusion	
12:00 pm	Lunch (Boxed Lunch To Be Provided)	
1:00 pm	ASK THE EXPERTS: DEEP ENDOVENOUS PROCEDURES Moderator: Peter Neglen, MD	
	Educational Objectives:	
	1.To be aware of technical aspects of venous stenting	
	2.To realize it role in chronic venous disease	
	3.To recognize its adjuvant role in early clot removal	
	Panelists: Haraldur Bjarnason, MD David Gillespie, MD Anthony Gasparis, MD	
2.00 pm	Coffee Break	

3:00 pm Coffee Break

SCIENTIFIC SESSION I: Endovenous

Moderators: Mark Meissner, MD Lowell Kabnick, MD

Educational Objectives:

- 1. Upon completion of this session attendees will understand:
- 2.An in depth analysis of endovenous ablative therapies comparing costs with open procedures.
- 3. Evaluating quality of life and disease severity outcomes when treating the superficial system with concomitant deep system reflux.
- 4. The safety of endovenous ablative procedures in patients that have or have had deep venous thrombosis.
- 5. Deciding if there is a benefit in performing immediate or staged adjunctive phlebectomy with endovenous ablation.
- 6. The importance of inflammatory markers in venous ulcer, and the affect of compression therapy on changes in cytokines levels and healing.
- 7. The importance of foam sclerotherapy in treating unusual varices of the sciatic nerve utilizing Duplex ultrasound guidance.

3:30 pm 1

1. Radiofrequency Ablation Versus Conventional Surgery For Varicose Veins - A Comparison of Costs

S. Subramonia¹, T. Lees² - ¹Queen's Medical Centre, Nottingham, United Kingdom, ²Freeman Hospital, Newcastle upon Tyne, United Kingdom

Background: To compare the costs involved (from procedure to recovery) following radiofrequency ablation and conventional surgery for isolated long saphenous vein incompetence in a selected population through a single-centre prospective randomised controlled trial.

Methods: Patients suitable for ablation (CEAP class 2-6) were selected by clinical assessment and duplex ultrasonography and randomised to either ablation or surgery and followed up one week and five weeks after surgery. Duplex ultrasonography was performed at first follow up. The primary outcome was the time (days) to return to full level of normal household activities. The hospital, practice and patient costs were calculated to indicate mean cost per patient under each category. Any difference in productivity between the two treatments in the employed group, due to sickness leave after surgery, was expressed as cost per working hour gained. The difference in productive unpaid household work after surgery between the two groups was expressed as cost per hour of household work gained.

Results: 93 patients were randomised. 88 patients (47-ablation, 41surgery) underwent the chosen intervention (five not treated). There was no follow up loss. Ablation took longer to perform than surgery (mean 76.8 vs 47.0 minutes, p<0.001, t-test). Patients returned to their normal activities (mean 5.87 vs 13.8days, p<0.001, t-test) and to work (mean 12.2 vs 19.8days, p0.006, t-test) significantly quicker following ablation than after surgery. Ablation was more expensive (£1275.90 vs £559.13 per patient) but produced a mean gain of one working week in the employed group compared to surgery. Based on 2005 Annual Survey of Hours and Earnings data (Office of National Statistics, UK) for full time employees, the cost per working hour gained was £6.94 (95% confidence interval 6.26, 7.62). Of those in the trial who were unemployed (20 patients) patients who underwent ablation returned to the normal household activities 17 days guicker than those who underwent surgery. Based on October 2000 Household Satellite Account data for household activities (Office of National Statistics, UK), the cost per hour of household work gained was £5.60 (95%) confidence interval 5.21, 5.99).

Conclusions: The increased cost of radiofrequency ablation is partly offset by a quicker return to work in the employed group. Reduced equipment costs and faster ablation with the new 'segmental ablation' catheter is likely to reduce the cost difference in future. Ablation is unlikely to be a cheaper option in the unemployed at reference year (2005) estimates.

 $\pounds = UK$ pound sterling

2. Endovenous Laser Ablation Improves Venous Outcomes Irrespective of the Presence of Deep Venous Insufficiency

B. S. Knipp, F. Mansoor, M. Hong, J. Bloom, E. Fellows, S. Blackburn, G. Adams, J. Pfeifer, D. Williams, T. Wakefield -University of Michigan, Ann Arbor, Ml

Background: We hypothesize that endovenous laser ablation therapy (EVLT) is equally successful in improving venous insufficiency in patients with or without deep venous insufficiency (DVI).

Methods: From January 2005 through August 2007, EVLT was attempted in 386 patients and 493 limbs. It was performed successfully in all but 17 cases. EVLT was performed alone in 334 limbs (70.2%) and with phlebectomy and/ or perforator ligation (EVLTP) in 142 limbs (29.8%). In a subset of patients undergoing EVLTP, perioperative thrombosis prophylaxis was administered based on a risk stratification protocol.

Results: Successful performance of EVLT led to complete saphenous vein ablation in 100% at one month and 90.3% at 1 year. Performance of EVLTP was associated with improved occlusion duration (P=.019). Mean revised VCSS (VCSS minus compression) preoperatively was 5.0 ± 2.9 , decreasing to 1.7 ± 1.5 beyond 360 days (P<0.001). Male gender (P=0.025) and performance of phlebectomy (P=0.020) were independently associated with greater improvement in scores with time. Changes in VCSS scores were equivalent regardless of DVI for both isolated EVLTP (Figures).

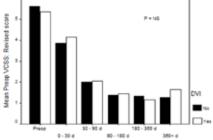
In cases of isolated EVLT, the true DVT rate was 0.6%, whereas for EVLTP, the rate was 2.8% (P=0.068); the rate of thrombus extension at the SFJ was 7.5% for isolated EVLT versus 2.8% for EVLTP (P=0.059). The use of risk-adjusted heparin prophylaxis in patients undergoing EVLTP did not have a significant effect on thrombotic complications. There were no differences in true DVT, thrombus extension, or superficial thrombophlebitis between patients with or without DVI (Table).

Using a multivariate regression model, obstructive pathophysiology was predictive of technical failure (OR 15.3, CI 2.3 to 100.1, P=0.004). Using a Cox proportional hazards model, performance of phlebectomy was independently predictive of duration of vessel occlusion (OR 8.3, CI 1.0 to 66. 7, P=0.047).

Conclusions: EVLT produces successful ablation and is associated with significant long-term improvement in VCSS. While there is a trend towards increased deep venous thrombosis when performing concomitant phlebectomy, there is also a significant improvement in long-term outcomes and VCSS scores, independent of DVI. Finally, the use of a risk-adjusted thrombosis prevention protocol had no effect on the rate of thrombotic complications from EVLTP.

Wednesday

Change in VCSS vs Time EVLT Only . Mean Preop VCSS: Revised score 5 P = NS 4 а. DVI 2 No ____Yes ٩, 30 90 el 360 6 0 - 30 d 90 - 180 d 300+ d Change in VCSS vs Time EVLT with Philebedomy e P = NS 5 4 3 2.



Thrombotic Complications					
	EVLT	EVLTP			
	DVI	No DVI	DVI	No DVI	
True DVT	2 (0.9%)	0 (0%)	4 (4.0%)	0 (0%)	
Thrombus Extension	17 (7.4%)	7 (7.6%)	4 (4.0%)	0 (0%)	
Thrombophlebitis	4 (1.7%)	1 (1.1%)	4 (4.0%)	2 (5.0%)	

4:10 pm

3.

Radiofrequency Ablation of the Great Saphenous Vein In Patients With Previous Venous Thrombosis: Is It Safe?

A. Puggioni, N. Marks, A. Hingorani, A. Shiferson, E. Ascher - Maimonides Medical Center, Brooklyn, NY

Background: The safety of radiofrequency ablation (RFA) of the great saphenous vein in patients with previous history of deep venous thrombosis (DVT) has not yet been determined.

Methods: From April 2003 to June 2006 we performed 293 consecutive RFA procedures in 274 patients (68% females) with a mean age of 60 ± 15 years. Patients with a history of previous venous thromboembolic events or with duplex scan evidence of post-thrombotic venous disease were identified in 29 of these cases (10%). These were compared to the remaining 264 cases (90%) without history or duplex evidence of thrombotic disease. Routine postoperative duplex scanning was obtained in all cases. Acute thrombotic (AT) events that developed in either lower extremity were documented and analyzed. According to the CEAP classification, 204 limbs (70%) were C2-C4, and 89 (30%) were C5-C6. Concomitant procedures included avulsion phlebectomy in 88 limbs (30%) and perforator surgery in 4 (1%).

Results: AT events after RFA were detected in 38 cases (13%) and included thrombus protrusion into the SFJ in 24 (8%), common femoral vein in 7 (2%) and calf vein DVT in 7 (2%). Overall incidence of AT in patients with and without evidence of previous DVT was 7% (2/29) and 14% (36/264), respectively (p=0.36). Advanced clinical presentation by CEAP classification(classes C5-C6) was not significantly different between the AT (15/38=39%) and non-AT (74/255= 29%) groups (p=0.19). Conversely, concomitant venous operations were associated with a significant increase in AT events (20/88= 23% versus 18/205=9%) with a p<0.002. All AT patients were successfully treated with standard anticoagulation. No cases of pulmonary embolism occurred in this series.

Conclusions: This experience shows that RFA of the great saphenous vein in patients with previous venous thromboembolic events is safe and should be offered as an alternative to surgical procedures. These data call attention to an increased incidence of AT events when concomitant venous operations are performed.

4:30 pm 4. Mid-Term Results of the Surgical Treatment of Varices By Phlebectomy With Conservation of A Refluxing Saphenous Vein

> P. Pittaluga¹, S. Chastanet¹, J. J. Guex² - ¹Riviera Veine Institut, Nice, France; ²Cabinet de Medecine Vasculaire, Nice, France

Background: A new physiopathological concept within superficial venous insufficiency (SVI) describes ascending progression from the collaterals to the saphenous veins (SV), leading to a treatment that aims to remove the varicose reservoir (VR) and not the SV. This study reports the mid-term results of this therapeutic approach.

Methods: This is a retrospective study of patients treated for varices by phlebectomy with conservation of the SV before July 2004. We evaluated the VR by determining the number of zones treated (NZT); each lower limb (LL) was divided into 32 zones. We performed a clinical examination and echo-Doppler after 6 months and 1 year, and then once a year until the 4th year. We monitored the progression of the saphenous reflux, as well as the signs and symptoms. We looked for risk factors (RF) for the persistence of SV reflux, an absence of clinical improvement, and postoperative varices recurrence.

Results: We operated on a total of 303 LL involving 221 patients (55 men and 166 women), with a mean age of 52.7 (20 to 93). All LL operated on presented preoperative SV reflux over 0.5 s: great saphenous vein (GSV) 85.8%; small saphenous vein (SSV) 11.9%; and GSV+SSV 2.3%. The NZT was 6.05 on average (2 to 10).

Saphenous reflux was abolished in 67.8%, 68.1%, 66.3%, 67.2% and 67.7% of cases respectively after 6 months, 1, 2, 3 and 4 years of followup. Symptoms improved or disappeared in 84.4%, 82.3%, 83.9% and 89.2% of cases, and cosmetic benefits were noted in 91.9%, 91.1%, 90.7% and 91.9% of cases at each annual check-up until the 4th year. The recurrence rate at 1, 2, 3 and 4 years was respectively 1.2%, 5.9%, 10.9% and 19.4%.

RF for SV reflux persistence were the preoperative existence of SV reflux reaching the malleolus and an NZT > 7.

Conclusions: Ablation of the VR with conservation of a refluxing SV is an effective treatment in the mid-term for the signs and symptoms of SVI, and leads to the abolition of SV reflux in over 2 out of 3 cases. The extent of the VR ablation is the key factor determining the hemodynamic and clinical efficacy of this conservative surgical approach.

5. Endovenous Laser Therapy With Concomitant Or Sequential Phlebectomy: A Randomized Controlled Trial A. I. Mekako, J. Hatfield, M. N. Abdul Rahman, S. Gulati, P. T.

McCollum, I. C. Chetter - Hull Royal Infirmary/University of Hull, Hull, United Kingdom

Background: Significant proportions of patients require secondary procedures such as sclerotherapy or phlebectomy following endovenous laser ablation of varicose veins. We compared endovenous laser therapy and concomitant phlebectomy (EVLTAP) with endovenous laser therapy (EVLT) only.

Methods: Patients undergoing EVLT were randomised to undergo concomitant phlebectomy (n=18), or no phlebectomy (n=18), and followed up at 1, 6, and 12 weeks post-procedure. Procedure duration, pain scores, return to work /normal activities, patient satisfaction, quality of life (QoL) outcomes, venous clinical severity scores (VCSS), and need for secondary intervention were compared. Results are expressed as median (inter-quartile range); p value.

Results: Duration of EVLTAP procedure was significantly longer than EVLT only: 67 (51-78) minutes versus 46 (38-56) minutes; p=0.003. There were no differences between groups in pain scores, time to work /normal activities, and patient satisfaction. EVLTAP patients had significantly lower Aberdeen Varicose Vein Scores at 6 weeks [7.12 (2.00-11.56) versus 14.74 (10.54-18.07); p=0.001] and 12 weeks [2.06 (0.00-6.71) versus 9.60 (7.08-13.39); p=0.009]. There were no significant differences between groups in any SF-36 domain at any time point. VCSS was significantly better in the EVLTAP group at 12 weeks. 6 patients (35%) in the EVLT only group required sequential phlebectomy, while no patient in the EVLTAP group required secondary procedures.

Conclusions: EVLTAP results in significantly better clinical improvement and disease-specific QoL outcomes than EVLT only, in the short-term. Although the procedure duration is longer, it neither increases pain nor delays return to work, and it obviates the short-term need for secondary procedures.

MINI PRESENTATIONS

5:10 pm

6.

(Mini Presentation 1) Foam Sclerotherapy of Venous Malformations V. Cheng – San Diego Vein Institute, Encinitas, CA

Background: Venous malformations are common and may occur either as localized or segmental lesions. They account for up to 80% of lesions in malformation clinics. They enlarge when dependent and with exercise. Skin temperature is usually normal and pain is variable but fairly common. Pathologically, the lesions are made up of anomalous dilated veins with irregularly thickened walls. Sometimes, interconnected channels penetrate normal tissues.

Radiologic imaging defines the extent of involvement of venous malformations, but 3D-MR venography is the best single imaging modality. It gives a bright hyper signal on T2-weighted spin-echo sequences and allows 3-D reconstruction. Because the lesions are usually low flow, Doppler ultrasound is useful as a preliminary imaging study and to monitor treatment.

The management of venous malformations will depend on the patient's age, cosmetic severity, and location of the abnormality.

Methods: During a 30 month period, 1427 patients were investigated at our institution for venous disorders and 1.2% (17 patients) were found to have venous malformations. The age range was from 15 to 76 years (mean 30.8±18.6). Nine patients had manifestations of lower extremity Klippel-Trenaunay (K-T) syndrome; seven had only venous malformations.

Foam polidocanol (2 or 3% concentration) was produced by the Tessari technique and duplex doppler was used for ultrasound guidance and to monitor effects of treatment.

Results: A goal was set for each patient. The mean number of treatments was 3.6 ± 2.8 (range 1-10). Pain free healing was achieved in patients with non-healing ulcerations and, cosmetically, all of the patients were improved. There was no need for anesthesia, analgesia, or hospital stay. No patient had a deep venous thrombosis.

Conclusions: Sclerosant foam is useful in treating low flow venous malformations. It is quick, painless, and effective while not requiring anesthesia or hospitalization.

7. (Mini Presentation 2) The Echo-Guided Sclerotherapy In Sciatic Nerve Varices Treatment

S. Gianesini, G. Tacconi, A. Palazzo, P. Fortini, E. Righi, E. Menegatti, A. Liboni, P. Zamboni - Ferrara University, Ferrara, Italy

Background: Sciatic veins are tiny deep venous plexuses that remain following embryologic involution of the primordial venous vessels, developing in the embryo alongside the nerves. These veins can be found dilated at the duplex scanning whenever a collateral venous drainage is required following a deep vein thrombosis or whenever we observe an angio-dysplasic disease afflicting the same vessels, and are defined sciatic nerve varices (SNV).

SNV are a possible cause of sciatic pain and they have to be considered in the differential diagnosis whenever the physical examination reveals the co-existence of varicose veins of the postero-lateral aspect of the leg.

Anatomically the SNV lay within the sciatic nerve and are composed of multiple channels spiraling around the same nerve. This data suggests not to proceed with a surgical approach in the treatment of these disease because of the obvious great risk of damaging the nerve structures.

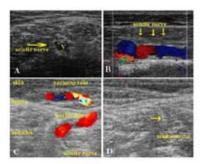
An alternative approach that we have considered in our study is the echoguided sclerotherapy.

The aims of this study is to verify both safety and effectiveness of ultrasonic guided foam sclero-therapy in treating SNV.

Methods: 19 consecutive patients, affected by symptomatic and angiodysplasic SNV, underwent both clinical and color Doppler investigation. All of them have been treated by one-shot echo-guided Tessari-foam sclero-therapy with just 2 of them who needed a second treatment always by the same technique. It has been possible to determine a follow-up for 12 of the 19 patients, that has included clinical as well as ultra-sonographic evaluation. Clinical results were assessed by objective and subjective Hobbs Scale. Objective evaluation was made by an independent assessor, who wasn't previously involved in the treatment. In addition duplex investigation was repeated. Subjective assessment was made by the patients at their own control.

Results: The mean follow-up lasted 24 months. Excluding skin pigmentation in 2 patients or a mild hardening feeling along the treated region in 3 other patients, nor minor nor major complications have emerged and the patients' compliance has been optimal. Reflux through the SNV, as the connected varicose veins, disappeared in the entire cohort. Mean subjective Hobbs Scale was A. Coexisting sciatic pain disappeared.

Conclusions: Ultrasonic foam sclero-therapy, in the mid term, seems to be both safe and effective, so representing a reliable and minimally invasive treatment of SNV.



A-B: Transversal (A) and longitudinal (B) high resolution B mode imaging of SNV.

C: Varices of the lateral aspect of the leg fed by reflux through a perforating vein by SNV.

D: The echo coming from the foam reaching its target in an area not so easily reachable by surgery.

6:00 pm Welcome Recepition

THURSDAY, FEBRUARY 21, 2008

7:00 am	Continental Breakfast / Exhibits Open
8:00 am	SCIENTIFIC SESSION II: Basic Moderators: Joseph Raffetto, MD & David L. Gillespie, MD
	Educational Objectives:
	!. Upon completion of this session attendees will gain knowledge in:
	 The importance of microparticles in the formation of venous thrombosis.
	3.The influence of matrix metalloproteinases (MMPs) in the venous ulcer wound environment, and how the composition of MMPs may affect venous ulcer wound healing.
	 The importance of physiologic age on thrombus formation and resolution.
	5. How MMPs affect venous relaxation by altering the calcium entry into smooth muscle.
	6. Important novel hemodynamic measurements in defining chronic venous disease.
	7.Neovascularization and a novel treatment algorithm.
	 Determining if cryo-venous stripping has any advantages over traditional stripping of the great saphenous vein.
8:00 am 8.	Microparticles Surface Proteins Influence Venous Thrombogenesis
	N. M. Abdullah M. Kachman A. Walker A. F. Hawley, S. K.

N. M. Abdullah, M. Kachman, A. Walker, A. E. Hawley, S. K. Wrobleski, D. D. Myers, Jr., J. R. Strahler, P. C. Andrews, P. K. Henke, T. W. Wakefield - University of Michigan, Ann Arbor, MI

Background: Microparticles (MPs) are small membrane vesicles released from a variety of cells upon activation. Elevated levels of MPs have been associated with many pathological conditions, including thrombosis and inflammation. MPs contain a unique subset of surface proteins derived from parent cell and may be responsible for their diverse biological functions. We used a quantitative proteomic approach to characterize the proteins which become amplified on the MP surface during venous thrombosis in an animal model.

Methods: Juvenile baboons (n4) underwent iliac vein thrombosis with temporary six-hour balloon occlusion as previously described. Plasma samples were taken at baseline and at 2 days post thrombosis for MPs analysis. MPs were extracted from platelet-poor plasma, digested separately with trypsin and tagged using isobaric tag for relative and absolute quantification (iTRAQ) reagents. The digests were subjected to 2dimensional liquid chromatographic separation followed by MALDI tandem mass spectrometry. Peak lists were generated and searched against all primate sequences using Mascot search engine. For protein identity, a minimum of two peptides at 95% confidence was required. Later, iTRAQ ratios were generated comparing relative protein level of day 2 to baseline. The proteomic analysis was performed twice for each blood sample, totaling to 8 experiments. Twenty-two proteins were considered to be on the surface of microparticles as determined by their consistent appearance in at least 4 out of 8 experiments. Data were normalized based on proteins (n8) that did not change their iTRAQ ratios at 2 days post thrombosis. Proteins were considered elevated or depressed if the iTRAQ ratio deviated by 20% change from normal value (1).

Results: Six proteins were statistically elevated with 1 protein being significantly depressed on day 2 versus baseline (Table 1). 15 proteins were present with no statistical elevation.

Conclusions: In this study we defined a diverse component of proteins that were associated with circulating microparticles in our animal model of venous thrombosis. These proteins influence thrombosis and inflammation through hemostatic plug formation (fibrinogen), stabilizing PAI-1 (vitronectin), inhibiting fibrinolysis (serpin peptidase inhibitor), stimulating antioxidant activity (haptoglobulin), and immunoregulation (Mamu IgM). Proteins not statistically elevated may play an important role during acute thrombosis as well. Proteomic data from this and future studies can be used to identify novel biomarkers to target future threapies for venous thrombosis. This study is the first to demonstrate the proteome of microparticles in a large animal model of venous thrombosis.

Table 1		
Protein Name	Average iTRAQ ratio	p-value
Fibrinogen gamma chain (elevated)	1.915	p<0.01
Fibrinogen beta chain isoform 4 (elevated)	2.001	p<0.01
Serpin peptidase inhibitor (elevated)	3.318	p<0.01
Vitronectin (elevated)	1.267	p<0.01
Haptoglobulin (elevated)	1.880	p<0.05
Trypsin precursor (elevated)	2.043	p<0.05
Mamu IgM-rh heavy chain (depressed)	0.750	p<0.01

8 :20 am

9. Inflammatory Cytokine Levels In Chronic Venous Insufficiency Ulcers Before and After Compression Therapy

S. Beidler, C. Douillet, D. Berndt, P. Rich, W. Marston -University of North Carolina at Chapel Hill, Chapel Hill, NC

Introduction: Patients with ulceration related to chronic venous insufficiency (CVI) have been reported to express high levels of proinflammatory cytokines from ulcer tissue. It is theorized that control of the inflammatory process is required for expeditious ulcer healing. In this study we evaluated the expression of pro-inflammatory cytokines and the multi-factorial cytokine TGF- β 1 in healthy and ulcer tissue before and after 4 weeks of high strength multilayered compression bandage therapy.

Methods: Tissue biopsies were obtained from ulcers in 30 patients with duplex confirmed venous insufficiency before and after four weeks of sustained high compression therapy. Healthy biopsies were taken from the ipsilateral thigh in 23 of the 30 patients. The tissue was homogenized and IL-1 α , IL-1 β , IL-6, IL-8, MCP-1, IFN- γ , MIP-1 α , MIP-1 β and TNF- α protein levels were obtained using a Luminex xMAP multiplexed assay which simultaneously measured cytokines in individual samples. TGF- β 1 levels were assessed by ELISA (N=10). All cytokines were normalized to total protein levels. Results were analyzed using ANOVAs, and data are presented as means and standard errors of the means.

Results: The average wound size decreased by 52% after 4 weeks of compression. Compared to healthy tissue, all pro-inflammatory cytokines, except IL-1 α , were significantly elevated in CVI ulcers before therapy. IL-6, MCP-1 and IFN- γ were elevated in the ulcer following therapy compared to healthy tissue (See Table 1). Compression therapy significantly decreased IL-1 β , IL-6, IL-8, IFN- γ and TNF- α ulcer tissue levels. IL-1 α healthy tissue samples were elevated compared to healthy tissue and continued to rise with compression therapy.

Conclusions: The majority of pro-inflammatory cytokines were significantly elevated in CVI ulcers when compared to healthy tissue using a multiplexed assay technique. Although all cytokines were identified in the ulcer tissue, disproportionately high levels of IL-8 were observed. Four weeks of compression therapy decreased the ulcer pro-inflammatory cytokine milieu and significantly increased TGF β -1 levels. The results indicate that the beneficial effect of compression therapy is mediated by a reduction of inflammatory mediators combined with an increased presence of TGF β -1.

Table 1. Pro-Inflammatory Cytokine and TGF- β 1 Protein Levels in Chronic Venous Insufficiency Ulcers Before and After 4 weeks of Compression Therapy.

Healthy Tissue			
(pg/ug protein)	Ulcer Before Therapy		
(pg/ug protein)	Ulcer After Therapy		
(pg/ug protein)			
IL-1α *^	2.639 + 0.637	0.88 + 0.313	0.29 + 0.043
IL-1β *#	0	0.163 + 0.051	0.023 + 0.01
IL-6 *^#	0	1.273 + 0.309	0.623 + 0.146
IL-8 *#	0.013 + 0.007	15.19 + 4	3.8 + 0.874
MCP-1 *^	0.108 + 0.038	1.028 + 0.1	0.937 + 0.106
IFN-γ *^#	0.010 + 0.004	0.275 + 0.049	0.139 + 0.029
TNF-α *#	0.003 + 0.001	0.019 + 0.005	0.010 + 0.001
MIP-1α*	0.048 + 0.014	0.542 + 0.186	0.244 + 0.028
MIP-1β *	0.113 + 0.013	0.316 + 0.092	0.191 + 0.019
TGF-β1 *^#	0.085 + 0.011	0.243 + 0.023	0.336 + 0.039

p < 0.05:

* Healthy Tissue vs. Ulcer Before Therapy

^ Healthy Tissue vs. Ulcer After Therapy

Ulcer Before Therapy vs. Ulcer After Therapy

8:40 am

10. The Matrix Metalloproteinase (MMP) Profile In the Venous Ulcer Bed May Provide A Prognostic Indication of Ulcer Healing

J. Tan, A. Smith, K. Burnand - Academic department of Surgery, Cardiovascular Division, London, United Kingdom

Background: Chronic venous ulcers are characterised by an imbalance in extracellular proteolytic activity causing abnormal collagen turnover. The aim of the study was to examine the activity of matrix metalloproteinases (MMPs) and their inhibitor TIMP1 (which are important regulators of extracellular matrix turnover) in chronic venous ulcer biopsies. The inhibitory effect of treatment with doxycycline on MMPs was also investigated.

Methods: Punch biopsies were taken from the ulcer margins of 55 patients with chronic venous ulceration of at least 6 months duration. Patients were randomised to receive doxycycline (200mg once a day orally) or placebo for 3 months. All patients were treated with 3-layer compression bandages and were followed up for a minimum of 12 months. A further punch biopsy was performed if the ulcer failed to heal. The activity and protein levels of MMP-1, -2, -3, -8, -9 and TIMP-1 were measured by bioimmunoassay and ELISA respectively.

Results: Thirty one patients achieved complete healing by 3 months. Significantly higher activities of MMP-2 (P= 0.0008) and MMP-8 (P= 0.0004) were found in those patients whose ulcer had failed to heal. Slow healers demonstrated a reduction in MMP-3 activity (P= 0.028). There was no difference in the total protein levels in all the MMPs studied between the two groups. Significantly higher levels of TIMP-1 (P=0.001) were found in the slow healers. Doxycycline suppressed the activity of MMP-1,-8 and -9 (P=0.025, 0.027 and 0.037 respectively). The administration of doxycycline did not appear to influence the leg ulcer healing rate.

Conclusions: There is a complex temporal pattern in MMP expression during ulcer healing, but it appears that increased MMP-2 and MMP-8 activities may inhibit ulcer healing. Selective pharmacological inhibition of these MMPs may be beneficial.

11. The Prothrombotic Effects of Aging On Acute Venous Thrombosis In A Rodent Model

A. P. McDonald, T. R. Meier, A. E. Hawley, J. N. Thibert, D. M. Farris, S. K. Wrobleski, P. K. Henke, T. W. Wakefield, D. D. Myers, Jr. - University of Michigan, Ann Arbor, MI

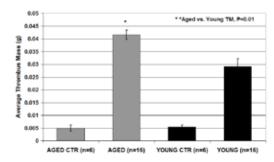
Background: Deep venous thrombosis (DVT) is recognized as a national health care concern. Recent studies have shown that the incidence of venous thrombosis significantly increases in the elderly. The risk of thrombosis increases by 1000 fold when the very old are compared to younger individuals. This current study identifies several age-related factors that may account for this increased risk.

Methods: Anesthetized male mice (C57BL/6, n=44) underwent complete inferior vena cava occlusion to produce venous thrombosis. Experimental groups included young mice (2 mo) vs. aged mice (11 mo), and aged match control animals from the same background. Mice were euthanized two days post-thrombosis for tissue harvest and blood collection. The following parameters were assessed: thrombus mass (g/cm), vein wall inflammatory cell populations (cells per 5 high powered fields), vein wall protein levels by multiplex ELISA (pg/ml), plasma PAI-1 activity (ng/mg), and circulating plasma microparticles (MPs, per 200 μ l of platelet-poor plasma).

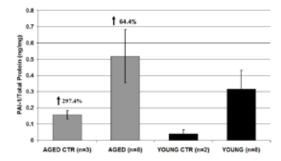
Results: Two days post thrombosis, aged mice had significantly larger thrombi than younger mice (Figure 1). Vein wall monocyte, lymphocyte, and total inflammatory cell populations were all significantly lower in aged mice (P<0.05). Vein wall protein analysis showed significant elevations in P-selectin at day 2 in aged mice, and significant declines in interleukin 4 (IL-4) and eotaxin protein levels versus young mice at the same time point (P<0.05). Aged control (CTR) mice had a 297.4% increase in PAI-1 activity compared to younger CTR animals. Two days post thrombosis, aged mice showed a 64.4% increase in PAI-1 activity compared to younger animals (Figure 2). The evaluation of mouse plasma showed aged mice to have a significant increase in circulating leukocyte-derived MPs compared to younger animals (5817±850 vs. 2563±283 MPs/200 µl PPP, P<0.01).

Conclusions: In our rodent model, aging significantly increased venous thrombosis. Aging significantly decreased vein wall inflammatory cell populations needed for thrombus resolution. Plasma PAI-1 activity increased with age suggesting PAI-1-dependent fibrinolysis is impaired in older mice. Both vein wall eotaxin and IL-4 protein levels, which influence fibroblast and endothelial cells, were significantly decreased with age. Aging significantly increased leukocyte-derived microparticle populations and procoagulant P-selectin activity that are known to promote venous thrombosis. This study shows that aging alters the balance of several prothrombotic and fibrinolytic factors to favor venous thrombosis.

Thrombus Mass



Plasma PAI-1 Activity



9:20 am 12. MMP-2 Induced Venous Relaxation Via Inhibition of Ca2+ Entry-Dependent Mechanisms of Venous Smooth Muscle Contraction

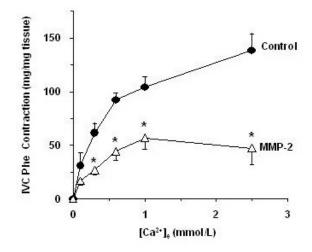
> J. D. Raffetto¹, R. Khalil² - ¹VA Boston Healthcare System, West Roxbury, MA, ²Brigham and Women's Hospital, Boston, MA

Background: Matrix metalloproteinases (MMPs) are implicated in the pathogenesis of varicose veins (VarV). We have shown that MMP-2 causes relaxation of venous segments and suggested a role of venous smooth muscle (VSM) hyperpolarization and activation of K+ channels; however, the downstream mechanisms involved are unclear. We tested whether MMP-2 induced venous relaxation involves inhibition of the Ca2+ mobilization mechanisms of VSM contraction.

Methods: Circular segments of rat inferior vena cava (IVC) were suspended between two wires in a tissue bath, and isometric contraction was measured. To test the role of Ca2+ release from the sarcoplasmic reticulum, IVC was incubated in Ca2+-free (2 mM EGTA) Krebs with or without MMP-2 (1µg/mL), then stimulated with phenylephrine (Phe, 10-5 M), caffeine (25 mM) or angiotensin II (AngII, 10-5 M). To test the role of Ca2+ entry through Ca2+ channels, after eliciting a transient Phe contraction in 0 Ca2+ Krebs, increasing concentrations of CaCl2 (0.1, 0.3, 0.6, 1, 2.5 mM) were added and the [Ca2+]e-contraction relation was constructed. Contraction data were presented as means±SEM mg/mg tissue.

Results: In IVC incubated in normal Krebs (2.5 mM Ca2+), Phe caused an initial peak (133±18) and a maintained contraction (73±12), while AngII caused a monophasic contraction (123±13). In Ca2+-free Krebs, caffeine did not cause contraction, suggesting limited role of the Ca2+induced Ca2+-release mechanism in IVC contraction. Also, in Ca2+ free Krebs, Phe and AngII caused a small contraction (7±2 and 15±3) that was not significantly affected by MMP-2 (10±3 and 73±27), suggesting little effect on IP3-induced Ca2+ release. The [Ca2+]e-contraction relation was reduced in MMP-2 treated IVC (figure), suggesting inhibition of Ca2+ entry. The specificity of the inhibitory effect of MMP-2 was demonstrated by its reversal with TIMP-1. In IVC treated with MMP-2, the Ca2+ channel blocker diltiazem (10 μ M) did not cause any further inhibition of Phe contraction, suggesting that Ca2+ entry is already inhibited by MMP-2.

Conclusion: In rat IVC, MMP-2 attenuates Ca2+ entry through Ca2+ channels, without affecting IP3-induced Ca2+ release from intracellular Ca2+ stores. MMP-2 role in venous dilation and varicose vein formation may involve inhibition of the Ca2+ entry mechanism of VSM contraction.



MINI PRESENTATIONS

9:40 am

13

(Mini Presentation 3)

Doppler Derived Maximum Venous Outflow Velocity (MVOV) Demonstrates Asymmetric Lower Extremity Venous Flow In Normal Individuals

M. Lebow, D. Cassada, O. Grandas, S. Stevens, M. Freeman, M. Goldman - UT Knoxville, Knoxville, TN

Background: It has been demonstrated in autopsy studies (33% of 107 cadavers) and more recently by the analysis of contrast CT scans (66% of 50 patients) that a significant portion of the population harbors asymptomatic compression of the left iliac vein by the right iliac artery. Venography lacks the sensitivity to be considered a true "gold standard" (66% sensitive) while IVUS is over 90% sensitive. No studies have addressed the physiologic difference in flow hemodynamics between the left and right lower extremities in normal individuals. We use Doppler ultrasound with induced high venous outflow to demonstrate asymmetric flow in the lower extremities of normal individuals.

Methods: MVOV in the common femoral veins were recorded using Doppler ultrasound on 30 volunteers. Inclusion criteria included females age 18-30 years, BMI <30, and no history of venous disease or leg swelling. All studies were performed by the same experienced vascular technologist. Volunteers were instructed to lay supine while a blood pressure cuff was inflated to 140mmhg around the mid-thigh. Presence of arterial flow was confirmed with color duplex after cuff inflation. Patients were instructed to exhale and hold their breath to augment venous outflow prior to rapid cuff release after 2 minutes. Outflow velocity and waveforms in the left and right common femoral veins were recorded for analysis.

Results: The mean age was 20.9 years (range 19-28 years) and the mean BMI was 21.9 (range, 18 - 25). MVOV was lower on the left side in 22 volunteers, lower on the right in 7 and equal in one. The mean right MVOV was 117.23 cm/sec (SD +/- 46.95) and the mean left MVOV was 95.44 cm/sec (SD +/- 32.94) (P= 0.01). The mean outflow acceleration on the left side was also lower (632 cm/s2) when compared to the right (961 cm/s2) (P=0.03) There was no correlation between left or right MVOV with age, BMI or height.

Conclusions: Significant differences in venous flow of the left and right lower extremities are present in normal individuals when venous outflow is increased over resting flow. This finding correlates with anatomic studies demonstrating a predilection towards narrowing of the left iliac vein in normal subjects. Doppler ultrasound is a simple, non-invasive method of quantifying venous hemodynamics that may aid in selecting patients for further diagnostic testing or intervention.

9:45 am 14 (Mini Presentation 4) Neovascularity and It's Treatment After Saphenous Ligation R. Bush, K. Hammond - Midwest Vein and Laser Center,

Dayton, OH

Background: Neovascularity is a leading cause for recurrent varices in those patients who have had a previous high ligation of the saphenous vein. Treatment of this condition remains somewhat controversial.

Methods: Thirty patients with neovascularity as an etiology of recurrent varicose formed the basis of this study. Ultrasound findings at the previous saphenofemoral junction (SFJ) and the outflow pattern that formed the recurrent varices was documented. All patients were followed for one year post treatment with foam sclerotherapy (Sotradecol - Bionesch) The number of treatments required as well as complications were noted.

Results: All patients had successful treatment of the neovascularity pattern at the groin level. Most patients require at least two treatments staged one month apart. This correlates with the number and size of channels present at the initial evaluation. The outflow pattern included flow into a retained saphenous vein, a duplicate saphenous vein, an accessory saphenous vein, or directly from the neovascularity channels into thigh varices. Ultrasound exam at one year revealed no recurrence of the neovascularity pattern. There was one case of partial femoral vein thrombosis as the only complication.

Conclusion: Neovascularity at the groin level after previous high ligation of the saphenous vein is a leading cause of recurrent varices. Foam sclerotherapy is a safe and effective therapy for neovascularity. Repeat groin operative procedures are contraindicated.

9:50 am 15 (Mini Presentation 5) Cryo Strip Versus Classic Strip of the Great Saphenous Vein

T. M. A. L. Klem¹, J. M. Schnater², P. R. Schutte³, A. C. van der Ham⁴, C. H. A. Wittens⁵ - ¹Erasmus Medical Center, Rotterdam, Netherlands; ²Academisch Medical Center, Amsterdam, Netherlands; ³Albert Schweitzer Hospital, Dordrecht, Netherlands; ⁴Sint Franciscus Hospital, Rotterdam, Netherlands; ⁵Haga Hospital, The Hague, Netherlands

Background: Stripping of the great saphenous vein (GSV) is one of the most performed operations in Europe. The classic strip with a plastic wire is the standard operation technique for stripping the GSV. A novel operation technique is the cryo strip, which uses a cryoprobe. The technique consists of a small incision in the groin, dividing the tributaries and retrograde insertion of the cryoprobe in the GSV till 5 cm under the knee. The cryoprobe then freezes to -85° Celsius (-121° Fahrenheit) after which the GSV is stripped by pulling the cryoprobe. We performed a randomized controlled trial to determine the efficacy of the cryo-strip versus the classic strip.

Methods: All patients with incompetence of the GSV and a patent deep venous system were included in the trial. Classic strip and cryo strip were performed in our outpatient clinic. All patients had a venous duplex preoperative and 6 months postoperative. Primary outcome was recurrence rate after 6 months of surgery confirmed by venous duplex.

Results: We included 539 patients and operated 494 patients.

Mean age was 51 year (20-84). There were 121 men and 373 women.

Cryo strip group: 249 patients, classic strip group: 245 patients.

Venous duplex after 6 months in the cryo-strip group:

An absent sapheno-femoral junction (SFJ) in 181 patients (73%), an incompetent SFJ in 49 patients (20%), an absent GSV in 173 patients (70%), an incompetent residual GSV in 54 patients (22%) and an occlusion of a residual GSV in 3 patients (1%). Missed to follow-up: 19 patients (8%)

Venous duplex after 6 months in the classic strip group:An absent SFJ in 175 patients (71%), an incompetent SFJ in 40 patients (16%), an absent GSV in 198 patients (81%), an incompetent residual GSV in 17 patients (7%). Missed to follow-up: 30 patients (12%)

Conclusions: The incidence of incompetence of the SFJ is higher in the cryo strip group than in the classic strip group. This difference is not significant.

The cryo-strip has a significant higher incidence of incompetence of the residual GSV (p<0.01)An explanation for this difference could be the difficulty of probe insertion in the GSV and perforation of the GSV. We think the cryo strip of the GSV is obsolete and should be abandoned.

SCIENTIFIC SESSION III: FOAM & DIAGNOSTICS

Moderator: Peter Pappas, MD

Educational Objectives:

- 1. Upon completion of this session attendees will understand:
- 2. If positioning of the patient undergoing foam sclerotherapy is important in preventing foam particle central migration.
- 3. The application of Duplex ultrasound and MRI in understanding the mechanism of pneumatic compression on the venous and muscles of the lower extremity.
- 4. The advantages or disadvantages in a randomized trial comparing treatment of varicose veins by surgery versus foam sclerotherapy.
- 5. The objectives of the National Venous Screening Program, how the NVSP has impacted the perception and education of the public and practitioners, and future direction in establishing nation-wide screening with emphasis on, risk of venous thromboembolism, venous clinical severity, primary care education, and timely specialty care referral and treatment.
- 6. The changes of MMPs and its naturally occurring inhibitors in the venous wall of varicose veins, and how this may impact on pathophysiology.
- 7.Demographic and risk factors that can effect pain following endovenous ablation.

10:30 am

16

Assessment of Techniques To Reduce Sclerosant Foam Migration During Ultrasound Guided Sclerotherapy D. A. Hill, R. Hamilton – The Vein Treatment Centre, Calgary, AB, Canada

Background: Endovenous chemical ablation is becoming an accepted technique for treatment of great saphenous vein insufficiency. However, echogenic phenomena in the right heart and high intensity transient signals detected by transcranial Doppler have been described by several investigators subsequent to foam sclerotherapy. An ischemic event following foam sclerotherapy of the great saphenous vein was reported recently in a patient with an occult patent foramen ovale. Concerns have also been raised about the effects of sclerosant foam on the pulmonary microvasculature. The present study compares the utility of 3 commonly used techniques for reducing sclerosant foam migration during ultrasound guided sclerotherapy of the great saphenous vein.

Methods: Group 1 consisted of 20 patients treated with ultrasound guided foam sclerotherapy of the great saphenous vein while lying supine with digital pressure used to occlude the saphenofemoral junction. In Group 2, 19 patients were injected with the leg elevated 30 degrees and digital pressure at the saphenofemoral junction. Group 3 involved 13 patients injected while the leg was elevated but without manual compression at the saphenofemoral junction. All patients were monitored with sub costal echocardiography during injection and for 3 minutes post injection.

Results: In Group 1, echogenic phenomena were demonstrated in the right heart in all 20 patients. In Group 2, echocardiography was positive

in 16 of 19 patients. In Group 3, echogenic cardiac phenomena were observed in 8 of 13 patients. There was a statistically significant difference in the incidence of echogenic phenomena between Groups 1 and Group 3 using Fisher's Exact Test (p=0.005). In Groups 1 and 2, a concentrated bolus of bubbles was frequently observed after release of digital pressure however only trace echogenic phenomena were seen in Group 3 where injection was performed with the leg elevated but no manual pressure at the saphenofemoral junction. No echogenic phenomena were observed in the left heart and there were no complications. Short term treatment results were equivalent among the 3 groups.

Conclusions: Endovenous chemical ablation of the great saphenous vein with foam sclerosants is best performed with the leg elevated and no digital pressure at the saphenofemoral junction. Further study is needed involving more subjects and foam produced from physiologic gases.

10:50 am

17 Combined MRI and Duplex Ultrasound Investigation of the Mechanism of Action of the Pneumatic Compression Devices

F. Lurie¹, H. Yoon², V. Scott³, R. L. Kistner¹ – 1University of Hawaii and Kistner Vein Clinic, Honolulu, HI, ²Hawaii Permanente Medical Group, Inc., Honolulu, HI, ³Keck School of Medicine USC, Los Angeles, CA

Background: The velocity and flow increase in deep veins is considered as the major mechanism DVT prevention by intermittent pneumatic compression (IPC). The origin of these hemodynamic changes has been never investigated by direct observation. It is unknown if the ICP displace the blood from small veins, superficial veins, or from compressed segments of deep veins. Although limited data on behavior of individual veins under compression is available, simultaneous measurements were never performed. The aim of this study was to measure changes in volume of subcutaneous tissue and veins, and subfascial tissue and veins caused by IPC, and to investigate their relationships to changes in blood flow in deep and superficial veins.

Methods: Five healthy volunteers with a mean age of 37 years participated in this study. The calf garments of two IPC devices, with different compression mechanisms were tested (WizAir -Medical Compression Systems, Inc, Ltd, Or-Akiva Israel, and VenaFlow -AirCast Inc., Summit, NJ).

The MRI was acquired with the garment pressure of 0 (baseline) and at maximum compression using a Philips 1.5T magnet (Philips Medical Systems, Bothell, WA) with a dedicated extremity coil. The following parameters were used: TR: 440 ms; TE 13 ms; 4 mm slice thickness with 0 mm gap; Field of view: 180 mm; Matrix 512x512 or 1024x1024; 20 slices were acquired; NEX: 4. Duplex scans were performed in same subjects to measure velocity and flow changes in the Great Saphenous and Femoral veins in lower thigh associated with using the two IPC devices.

Results: Compression with both of the tested devices was associated with measurable decrease in volume of subqutaneous tissue under the garment (p<0.001), total volume of superficial veins (p=0.004), and volume of the GSV (p=0.038). There were no measurable changes in subfascial volume of the calf under the garment.

Flow increase in FV and GSV under compression highly correlated with decrease in volume of superficial veins (r=0.77 and r=0.74 respectively), but not with changes in deep veins volume (r=0.3 and r=0.15). Increase in flow in FV was also highly correlated with increase in GSV flow (r=0.79). A single strongest predictor of venous flow increase was the change in subqutaneous veins volume (r square=0.73, p=0.0002 -linear regression).

Conclusions: 1. MRI imaging provides valuable information on changes in intra- and extra - venous volumes of the extremity under compression.

2. Displacement of blood from the subcutaneous veins into the deep venous system is most likely the major contributor to hemodynamic action of the IPC.

 11:10 am
 18 Comparison Between Surgical Treatment and Ultrasound-Guided Microfoam Sclerotherapy For Patients With Primary Varicose Veins In the Lower Limbs: Early Results of A Randomized Controlled Trial M. Figueiredo, S. P. Araujo, F. Miranda Jr - Escola Paulista de Medicing - Unifesp, Uberlandia, Brazil

BACKGROUND: This is a prospective study carried out by comparing patients with chronic venous insufficiency (CVI) under the C5EpAsPr CEAP classification submitted to operative treatment versus microfoam echoesclerotherapy over a period of six months.

METHODS: Sixty patients were selected with a number of them submitted to operative treatment with overall saphenectomy, removal of collaterals and perforating vein incompetent (n= 29) and microfoam echoesclerotherapy using the Tessari technique. The primary endpoints were the clínical scores of venous severity under the CEAP clínical classification (pain, oedema, inflammation, hyperpigmentation and lipodermatosclerosis); the secondary endpoints assessed effectiveness of the vascular ultrasound therapy and the complication arising from the treatments.

RESULTS: The two patient groups were compared both before and after treatment, and were shown to be statistically equal. The inflammation, hyperpigmentation and lipodermatosclerosis scores were also shown to have improved under both treatments employed (statistically not significant). In the assessement with the vascular ultrasound, the efficacy in the microfoam echosclerotherapy was 77.8% (n/N-21/27) against 89.6% (n-N 26/29) of the patients submitted to operative treatment in the final assessement of 180 days (statistically not significant).

CONCLUSIONS: Improved clinical scores in the CEAP venous severity (pain, edema, inflammation, hyperpigmentation and lipodermatosclerosis) in both surgical treatment and echosclerotherapy for period six month.

Pacientes	Echoesclerotherapy	Surgery
N- all patients	27	29
n- sucess	21	26
%n/N	77,8	89,7





Segment	Surgery	Surgery	Surgery	Echoesclero- therapy	Echoesclero- therapy	Echoesclero- therapy
	Treatment	Sucess	%	Treatment	Sucess	%
Great Saphenous vein (thigh)	27	26	96,30	25	20	80
Great Saphenous vein (below knee)	23	21	91,30	24	16	66,67
Short Saphenous vein	3	3	100	4	2	50
Collateral thigh	1	1	100	2	2	100
Collateral below knee	9	7	77,78	12	10	83,3
perforator thig	1	1	100			-
perforator below knee	8	7	87,50	7	4	57,1
Total	72	66	91,67	74	54	72,97

MINI PRESENTATIONS

(Mini Presentation 6)

11:30 am

19

National Venous Screening Program – An Update Marc Passman, MD

11:35 am **20 WITHDRAWN**

11:35 am **21** (Mini Presentation 8)

Patient Characteristics and Treatment Factors That Affect Pain Following Endovenous Laser Treatment (EVLT) For Venous Insufficiency

P. A. Hertzman, B. Peterson² - ¹Vein Care of New Mexico, Los Alamos, NM, ²University of New Mexico, Albuqueque, NM

Background: Pain following EVLT for venous insufficiency is common, yet limited information is available regarding what factors influence its incidence or severity. Our objective was to determine the effect of patient characteristics and treatment factors on pain following EVLT.

Methods: Incompetent veins were treated using ultrasound guidance with a 14 Watt, 810 nm diode laser after tumescent anesthesia. The laser fiber was withdrawn at 1-2 mm per second delivering between 75 and 140 joules per cm. Patients recorded pain on a 0-10 scale (0=no pain; 10=most severe pain) beginning on the day of treatment. All patients were seen and duplex ultrasound was performed one week following EVLT. We analyzed results for both peak pain and mean pain for the first seven days for 40 procedures in 27 patients (Females-17; Males-10; Mean Age=50.52 \pm 11.91 years; Range=28-81 years; Mean BMI= 27.23 \pm 5.93 kg/m2)by linear regression analysis, Pearson product-moment correlation coefficients, and t-test comparisons.

Results: The treated vessels had closed in all cases and no DVTs were identified. All patients experienced some pain during the first week which was mild on average (mean = 3.437 SD= 2.314; range= .29 - 10). Pain most commonly peaked on either Day 2 - 42.5% (n=17) or Day 7 - 45.0% (n=18). The Peak pain scores were mild to moderate (mean = 4.925; SD=2.615; range = 1-10). The peak use of analgesics was on Day 1. The most commonly used medications were Tylenol and Ibuprofen. No pain medications were used on 37% of the patient-days during the first week. The means, correlations (R) with pain, and P values for patient characteristics and treatment factors are shown in this table:

Peak Pain	Mean Pain							
Patient Characteristics	N		R	Р	R	Ρ		
Gender	27		0.397	0.011	0.330	0.037		
Leg treated: Right = 20; Left =20	40		0.126	0.439	0.078	0.631		
Vein GSV=28; SSV=9; other=3	40		0.425	0.006	0.324	0.041		
	N	Mean	SD	Range	R	Р	R	Р
Age (years)	27	59.52	11.91	28-81	0.105	0.521	0.052	0.748
BMI (kg/m2)	24	27.23	5.93	18.5- 38.6	0.091	0.596	0.137	0.427

Max Vein Diameter (mm)	30	8.27	3.32	4.1-17.9	0.444	0.014	0.451	0.012
Access vein diameter (mm)	40	4.97	1.34	1.59-8.7	0.274	0.150	0.285	0.134
Treatment Factors								
Laser Treatment Time(sec)	40	206.65	60.59	101-299	0.345	0.029	0.323	0.042
Vein Length treated (cm)	38	31.03	10.02	15-47	0.320	0.050	0.384	0.017
Laser Power total (joules)	40	2858.50	843.25	1414- 4185	0.303	0.057	0.263	0.101
Laser power (J/cm)	38	91.44	5.51	80.9- 110.7	0.126	0.450	0.114	0.497

T-test comparisons showed increased pain in women vs men (P<0.05)) and for GSV vs other veins(P<0.05)

Conclusions: Pain during the week following EVLT is on average relatively mild even at its peak and rarely requires pain medication stronger than over the counter analgesics. The level of pain is significantly more common in women, when treating the GSV, and in veins with larger maximum diameters. Treatment factors significantly associated with increased pain include laser treatment time and length of the vein treated.

12:00 pm	AMERICAN COLLEGE OF PHLEBOLOGY SCLEROTHERAPY SESSION Moderator: Steve Zimmet, MD & Nick Morrison, MD
	Educational Objectives:
	1.Better utilize sclerotherapy to treat incompetent varices
	2. Minimize risk of complications of sclerotherapyy
	3. Understand issues related to the importation, compounding and off-label use of sclerosants
12:00 pm	Office Set-Up and Sclerotherapy Techniques
	Nick Morrison, MD
12:15 pm	Sclerotherapy: Cleaning Up Before & After Endovenous Laser
	Robert Min, MD
12:30 pm	X-Ray Guided Sclerotherapy
	Mel Rosenblatt, MD
12:45 pm	Sclerosants: Importation, Compouding and Off- Label Use
	Steve Zimmet, MD
1:30-5:50 pm	INDUSTRY WORKSHOPS (Three 80-Minute Sessions)
	Ultrasound Investigations for Venous Disease Moderator: Nicos Labropoulos, MD
	Educational Objectives:
	 Understanding of basic normal venous anatomy identified by venous ultrasound.
	2. Understanding of diagnostic criteria for venous thrombosis using venous ultrasound.
	3. Understanding of diagnostic criteria for venous insufficiency (deep, superficial, perforator) using venous ultrasound
	Endovenous Ablation of the Saphenous Vein Moderator: Michael Vasquez, MD
	Educational Objectives:
	 Understand and perform U/S guided access of an enlarged saphenous vein based on practice on a model.
	2. Discuss the importance and technique of intra-compartmental tumescent anesthesia for the performance of endovenous saphenous vein ablation.
	3. Identify different modalities of saphenous vein ablation for possible integration into their practice.

Pharmaco-Mechanical Thrombectomy (PMT)

Moderator: Peter Lin, MD

Educational Objectives:

- 1. Understand endovascular treatment strategies of acute deep venous thrombosis.
- 2. Understand the role of mechanical thrombectomy in the treatment of acute deep venous thrombosis
- 3. Be familiar with various mechanical thrombectomy devices in the treatment of acute deep venous thrombosis.
- 4. Have insight into potential applications of pharmacomechanical thrombectomy in acute deep venous thrombosis.

Venous Ulcer Wound Care

Moderator: William Marston, MD

Educational Objectives:

- I.Evaluate various methods of compression and the advantages and disadvantages of each in the treatment of venous leg ulcers
- 2. Consider the vast range of products available to apply to the wound surface of venous leg ulcers and learn strategies to choose the best ones for each leg ulcer
- 3.Review the active therapies available that are proven to accelerate the healing of leg ulcers and demonstrate proper application techniques for these products

OR CONCURRENT SYMPOSIUM SESSION

1:30 – 2:50 pm	Venous Coding and Maximizing Reimbursement Moderator: Robert Zwolak, MD
	Educational Objectives: 1.Use appropriate category one CPT codes to report standard venous operations 2.Understand the requirements for development of new CPT codes 3.Have a working familiarity with the method by which CPT codes are valued
3:00 – 4:20 pm	Venographic Assessment Moderator: David Gillespie, MD
	Educational Objectives: 1.Understand the indications and techniques for performing ascending venography 2.Understand the indications and techniques for performing ilio/ cavography 3.Understand the indications and techniques for performing ovarian vein/pelvic dumping imaging
3:00 – 3:15 pm	Ascending/Decending Venography David Gillespie, MD
3:15 – 3:30 pm	Venographic Assessment of PelvicCongestion Syndrome Mark Meissner, MD
3:30 – 3:45 pm	Extremity Venography For Venous TOS Marc Passman, MD

What To Do With Recurrent Varicose Veins?

Moderator: Andre van Rij, MD

Educational Objectives:

- 1. Understand the causes of recurrence of varicose veins and the role that neovascularisation has in this
- 2. Gain a basic understanding of the biology of neovascularisation, and recanalisation,
- 3.Be aware of how this varies with different treatments of varicose veins and how it might be prevented.
- 4.Be familiar with treatments for recurrence and their relative merit.
- 5. Have a rationale for counseling patients regarding the risk of recurrence following varicose vein treatment.

PLEASE NOTE: The following evening symposium is included in the registration fee for physicians and allied health professionals. However, seating is limited and pre-registration is required. We regret that due to strict codes, spouses and guests may not attend.

6:30 - 8:30 pm EVENING SYMPOSIUM

Supported by Bacchus Vascular and Sanofi Aventis.

THE TIMES THEY ARE A-CHANGING: VENOUS THROMBOEMBOLISM UPDATE 2008

Educational Objectives:

- 1.Be familiar with the latest ACCP Chest Guidelines
- 2.Understand current concepts regarding the treatment choices and duration of treatment for venous thromboembolism
- 3. Understand the differences between iliofemoral thrombosis and other forms of venous thrombosis

6:30 – 6:50 pm	Putting New Joint Commission Quality Standards For
	DVT Into Hospital Practice
	Joseph Caprini, MD

6:50 - 7:10 pm The Role of IVC Filter Placement In DVT Prophylaxis and Treatment David Gillespie, MD

7:10 - 7:30 pm The Relationship Between the Location of Thrombosis and the Severity of the Post-Thrombotic Syndrome Peter Neglan, MD

- 7:30 7:50 pm Iliofemoral DVT Thrombus Removal Techniques: Safe and Effective Michael Zatina, MD
- 7:50-8:10 pm Mechanolytic Intervention For Iliofemoral DVT and the Need For A RCT Anthony Comerota, MD
- 8:10-8:30 pm Panel Discussion/Q&A

FRIDAY, FEBRUARY 22, 2008

7:00 am	Continental Breakfast / Exhibits Open
7:30 am	SCIENTIFIC SESSION IV: CHRONIC VENOUS DISEASE Moderators: Michael Ricci, MD & Marc Passman, MD
	Educational Objectives:
	1.Upon completion of this session attendees will be able to:
	Understand the variability in venous Duplex laboratory reporting and the need for uniformity.
	3. Understand the important application of intravascular ultrasound in placement of vena cava filters in multi-trauma patients with emphasis on technique, anatomic variations, complications, and early durability.
	4.A novel technique utilizing an endovenous valve stent for treating patients with deep venous insufficiency.
	 Understand the hemodynamic relation between the iliac venous system and the saphenofemoral junction.
	6.Appreciate the complex nature of varicose vein formation and chronic venous disease, and the role that hypoxia may have on its pathogenesis.
7:30 am	22 Use of A Structured Audit Tool In Assessing Venous Duplex Reports D. L. Wooster - University of Toronto, Toronto, ON, Canada

Background: Venous duplex reports of studies performed to investigate patients with leg pain and swelling (suggestive of deep venous thrombosis) have been noted to be of variable quality. A structured audit tool, based on published guidelines, can give quantitative assessments of such reports. Service gaps, as defined by technical protocol and performance gaps and physician interpretation gaps, can be identified in discrete areas of the duplex study by such an audit. A variety of focused interventions have been suggested to stimulate guideline implementation and continuing medical education around these issues. The aims of this study were to apply the audit tool to identify service gaps and use the results to identify strategies for quality improvement.

Methods: The venous component of a previously validated structured audit tool was applied to 100 venous duplex reports from community labs and hospital imaging departments. The audit tool contained 28 elements in 8 domains, based on SVU, ICAVL, CPSO and RSNA ultrasound practice standards. The audit allowed for quantitative scoring of each domain. The results were analyzed to identify common themes in order to infer areas for guideline implementation and focused education initiatives.

Results: The analysis of 100 reports from 7 labs and 4 hospitals showed overall average scores of 2.6 / 5 (range 1.8 - 4.2). Of the 8 domains, demographics (3.9), indication (3.9) and test performed (3.5) compared

favourably to other applications of the tool. Description of the test (2.2), findings (2.3), limitations (1.8), interpretation (2.8) and overall clinical applicability (2.6) fell below acceptable norms. Labs with ICAVL status performed better; there was no difference between community- and hospital-based facilities otherwise. Thematic analysis of the results showed a) systems issues, b) knowledge gaps, c) application gaps and d) overall service gaps in venous duplex study reports. Educational interventions were identified for each issue and repeat audits demonstrated practice change outcomes.

Conclusions: Multiple gaps are identified in venous duplex reports. Focused interventions are recommended to improve the quality and clinical relevance of such reports. Repeat audits of selected domains can be used to monitor and document quality improvement. :50 am

23 Intravascular Ultrasound Guided Inferior Vena Cava Filter Placement In the Multi-Trauma Patients From Global War On Terrorism: A Single Center Experience G. Aidinian, A. A. Amin, P. W. White, E. Adams, C. J. Fox, M. Cox, D. L. Gillespie - Walter Reed Army Medical Center,

Washington, DC

Background: Blast and high velocity fragments have resulted in a multitude of complex extremity and soft tissue injuries. Consequently, there is increased risk of venous thromboembolism from prolonged immobilization, inability to use extremity compression devices, and interrupted anticoagulation. In a percentage of these patients, intravascular ultrasound (IVUS) guided inferior vena cava (IVC) filter insertion was used due to renal failure, or high transport risk due to pulmonary failure, open abdomen, or hemodynamic instability. The objective of this study was to review our experience in bedside IVUS guided insertion of IVC filters in military multi-trauma patients.

Methods: A retrospective analysis was performed of all IVUS guided bedside IVC filter placed by the Vascular Surgery Service at Walter Reed Army Medical Center between August 2003 and October 2007. Abdominal x-rays were performed on all the patients to document filter location. Injury Severity Score (ISS) was calculated for each patient and compared to published civilian controls.

Results: Over the time period studied there were a total of 66 patients who underwent IVC filter placement. Fourteen of these patients had bedside IVC filter placed under IVUS. There were 13 males and 1 female with average age of 26 years. The mean (\pm SD) ISS for military bedside IVUS placement was 37.2 (\pm 9.9) compared to 25.1 (\pm 2.2), reported in civilian trauma patients (Rosenthal et al. J Vasc Surgery 2004;40:958-64). The most common cause of injury was from explosive devices (57%), followed by gun shot wounds (28%), rocket propelled grenade (7%), and motor vehicle crash (7%). Average hospital length of stay was 43 days. 13 patients had Günther Tulip filter inserted and 1 patient had TrapEase filter inserted. Indications for filter insertion were deep venous thrombosis (DVT) in 36% of patients, and pulmonary embolus (PE) in 28%. 35% had IVC filters inserted prophylactically.

Conclusion: Military trauma population ISS is considerably higher than what is reported in the civilian population. The bedside IVUS guided IVC filter insertion is particularly useful in this population with the renal failure, or high transport risk due to pulmonary failure, open abdomen, or hemodynamic instability.

78:10 am

24 The Effect of Pressure On Migration and Further Characterization of the Venous Ulcer Fibroblast

G. Scriver, A. Stanley, M. A. Ricci, K. Corrow, M. Slusarczyk, S. Shackford, J. Adams, G. Steinthorsson, D. Berges, A. Howard - University of Vermont, Burlington, VT

Background: Elevated atmospheric pressures have been found to alter in vitro fibroblast wound healing. An important aspect of wound healing involves the complex process of fibroblast migration. We hypothesized that neonatal fibroblasts (NNF) grown under elevated pressure will demonstrate delayed migratory ability and show an attenuated response to the mitogen PDGF. In addition, it has been hypothesized that fibroblasts subjected to years of chronic venous insufficiency (CVI) respond by aging prematurely and achieve a state of cellular senescence. If this is the case, and if pressure is the physiologic stimulus for the phenotypic changes and premature aging seen in both fibroblasts grown from venous ulcers and NNF grown under pressure, then these cell populations should both demonstrate shortened telomere lengths.

Methods: A pressurized incubator was used to culture neonatal fibroblasts at atmospheric pressure (ATM) and at ATM + 60 mmHg for 2 weeks. Cells were then placed onto a fibronectin coated permeable membrane. Cells were treated with PDGF- $\beta\beta$ or left untreated, were fixed after 12 or 24 hours, and the mean number of migrated cells were recorded under each experimental condition. The pressure incubator was also used to culture NNFs at ATM and ATM + 120 mmHg, as well as venous ulcer fibroblasts at ATM for a period of 2 weeks. DNA was isolated from these cells, and PCR was performed on the southern blots to determine telomere restriction fragment length among each cell population.

Results: Independent of PDGF- $\beta\beta$ application, fibroblasts grown under elevated pressure demonstrated less migration than NNFs grown at atmospheric pressure (P=0.004). Stimulation with PDGF- $\beta\beta$ increased the migration rate among both NNFs under elevated pressure and NNFs at atmospheric pressure, as compared to the untreated cells (P=<0.001). The migratory response to PDGF- $\beta\beta$ stimulation was not significantly different between NNFs grown under elevated pressure and NNFs at atmospheric pressure (P=0.17). Telomere length analysis demonstrated that the restriction fragment length was identical among NNFs under pressure, NNFs at atmospheric pressure, and venous ulcer fibroblasts.

Conclusions: This study demonstrated that elevated pressure alters fibroblast function, leading to a decreased in vitro migratory ability in these cells. However, NNFs grown under pressure retained their ability to migrate in response to PDGF- $\beta\beta$ stimulation. The finding of equivalent telomere lengths in venous ulcer fibroblasts, NNFs under pressure, and NNFs at atmospheric pressure suggests that while the effect of pressure on fibroblasts may induce cellular changes similar to premature aging, that these fibroblasts do not become truly senescent.

8:30 am

EUROPEAN VENOUS FORUM -FIRST PLACE WINNER

Haemodynamic Assessment of Iliac Veins and Their Relation With the Sapheno-Femoral Junction

P. Brazis¹, R. Piotrowicz¹, N. Labropoulos², A. Jawien¹ ¹Department of Surgery, Collegium Medicum, Nicolaus Copernicus University, Bydgoszcz, Poland - ²Division of Vascular Surgery, University of Medicine & Dentistry of New Jersey New Jersey Medical School, Newark, NJ USA

Purpose: Iliac vein reflux has not been evaluated in patients with chronic venous disease (CVD). This prospective study was designed to determine the prevalence of iliac vein reflux in relation to saphenofemoral junction (SFJ) reflux in the whole spectrum of CVD.

Methods: One hundred and forty three limbs in 72 patients were prospectively evaluated by duplex scanning and clinical examination. Patients were included from the whole spectrum of CVD and were graded according to the CEAP classification. The iliac veins were assessed just above the inguinal ligament in standing position by the Valsalva test. The cut-off value for iliac and femoral vein reflux was set at >1s and for the superficial veins at >0.5s. SFJ was examined at four points above and below the terminal and preterminal valves. The diameters of all the relevant veins were measured as well.

Results: Iliac reflux was found in 58 limbs (40.5%). The prevalence of iliac reflux was significantly higher in more advanced CEAP classes C4-C6 41(28.7%) vs 16(11.2%) in C1-C3, p<0.05. Iliac reflux was associated with SFJ incompetence. At this level reflux was detected more frequently in patients with iliac reflux than in patients without it 48(88.9%) vs 6(11.1%), p<0.05. The diameters in SFJ region were also significantly larger in patients with iliac reflux compared to those ones without (CFV 16.2 vs 15.1 and SFJ 7.5 vs 4.6 mm, p<0.05 for both).

Conclusion: Iliac vein reflux is significantly more prevalent in the presence of SFJ reflux and suggests that the presence of the latter is the cause of the former. The diameters of CFV and SFJ are greater in the presence of iliac vein reflux. These findings are more evident in patients with skin damage and provide a good hemodynamic explanation for the severity of the disease.

EUROPEAN VENOUS FORUM -SECOND PLACE WINNER

Is Hypoxia A Feature of Varicose Vein Disease?

B. Sharp^{1,2}, B. T. Navin¹, C. Monaco^{1,2}, E. Paleolog^{1,2}, A. H. Davies² - Kennedy Institute of Rheumatology, Imperial College, London, UK; Charing Cross Hospital Department of Surgery, Oncology and Anaesthetics (SORA), Imperial College, London, UK

Background: An imbalance between matrix metalloproteinases (MMP) and the MMP inhibitors TIMP is thought to play an important role in varicose vein disease through alterations in the extracellular matrix. It is likely that MMPs/TIMPs are stimulated by inflammatory cytokines. Hypoxia has been previously suggested to be involved in varicose vein disease. Hypoxic episodes are known to influence, through the modification of transcription factors, the release of cytokines and growth factors e.g. VEGF. The most important transcription factor is Hypoxia inducible factor (HIF). In this study we investigated whether the HIF family members HIF-1 α and HIF-2 α were present in varicose veins, and whether their presence influenced certain MMPs/TIMPs and cytokines that are already thought to play a role in varicose vein disease.

Materials And Methods: Varicose vein segments were obtained from 15 patients undergoing corrective surgery. RNA extraction was performed on the vein tissue and the message levels for MMPs (MMP-2, MT1-MMP), TIMPs (TIMP-2, TIMP-3), cytokines (VEGF, TNF- α , IL-1, IL-6), HIF-1 α and HIF-2 α were quantified by Sybr Green I PCR (Polymerase chain reaction), and expressed relative to a pool of mRNA from healthy vein.

Results: Both HIF-1 α and HIF-2 α were expressed in varicose veins. There was significantly lower expression of HIF-1 α (median=0.7950) than HIF-2 α (median=8.105) in proximal varicose vein segments (Mann-Whitney p=0.0010).

HIF-1 α mRNA showed significant correlation with MMP-2 (p=0.0211) and TIMP-2 (p=0.0435) mRNA levels. There was no significant relationship between HIF-1 α and either MT1-MMP or TIMP-3. When compared with cytokine expression there was a significant correlation of HIF-1 α with VEGF (p=0.0431), TNF- α (p=0.0347) and IL-1 α (p=0.0122). However, there was no correlation between HIF-1 α and IL-6.

HIF-2 α showed very significant correlation with MT1-MMP (p=0.0043). There was no correlation between HIF-2 α and MMP-2, TIMP-2 or TIMP-3. In terms of cytokine mRNA levels, HIF-2 α expression showed very significant correlation with TNF- α (p=0.0347) only.

Conclusion: For the very first time we have shown that HIF-1 α and HIF-2 α are expressed in varicose veins. HIF-2 α was significantly more abundant in proximal varicose veins. HIF-1 α displayed a statistically significant correlation with MMP-2 and TIMP-2. In contrast, HIF-2 α showed a very significant correlation with MT1-MMP. When correlating HIFs with the cytokines, HIF-1 α showed significant correlation with VEGF,

TNF- α and IL-1, whereas HIF-2 α showed very significant correlation with TNF- α . These data suggest a potential association between hypoxia and cytokines in regulating MMP/TIMP balance and thus altering the extracellular matrix in varicose vein disease.

9:10 am Coffee Break / Visit Exhibits

9:40 am	SCIENTIFIC SESSION V: ENDOVENOUS STENTING Moderators: Robert McLafferty, MD & Peter Neglen, MD
	Educational Objectives:
	1. Upon completion of this session attendees will be able to:
	Understand the variability in venous Duplex laboratory reporting and the need for uniformity.
	3.Understand the important application of intravascular ultrasound in placement of vena cava filters in multi-trauma patients with emphasis on technique, anatomic variations, complications, and early durability.
	4.A novel technique utilizing an endovenous valve stent for treating patients with deep venous insufficiency.
	 Understand the hemodynamic relation between the iliac venous system and the saphenofemoral junction.
	6.Appreciate the complex nature of varicose vein formation and chronic venous disease, and the role that hypoxia may have on its pathogenesis.

9:40 am 25 Venous Stenting Across the Inguinal Ligament P. Neglén, P. Tackett, S. Raju - River Oaks Hospital, Flowood, MS

Background: Stenting of arteries across joints is prohibited due to frequent stent fracture and secondary stenosis or occlusion. The aim of this study is to assess the fate of venous stents placed across the inguinal ligament.

Methods: During 1997 to 2005, 144 limbs had iliofemoral stenting performed extending beneath the inguinal ligament into the common femoral vein for non-thrombotic iliac vein lesions, so-called compression lesions (NIVL; 22 limbs), chronic thrombotic non-occlusive obstruction (PTS-obstr; 80 limbs), and chronic thrombotic occlusion (PTS-occl; 42 limbs). Female/male ratio = 107/37; left/right limb = 95/49; median age 56 years, range: 22-86; previous DVT 122 limbs. Braided stainless stents were most frequently used, but in 12% of limbs nitenol mesh stents were placed. The patients were followed with venography to assess patency and stent integrity 3 and 9 months post-intervention and then annually.

Results: The patients were followed for mean 24 months (range: 1 - 91 months) Twenty stent systems occluded during the observation period; 8 limbs had successful and 5 limbs failed removal of thrombus, 7 limbs had no intervention. Intervention of patent stents was performed in 24 limbs; 19 limbs had balloon angioplasty of in-stent restenosis (ISR), 3 had further distal stent extension of the CFV, and one had proximal extension into the IVC. Only one stent was crushed at the inguinal ligament, a nitenol stent, which was treated by insertion and angioplasty of a braided stainless steel stent. No other stents were compressed at this site and no stent fractured. The overall assisted-primary and secondary patency was at 42 months 81% and 88% (NIVL 100% and 100%; PTS-obstr 88% and 92%; PTS-occl 66% and 78%, respectively).

Conclusions: Contrarily to arterial stenting, venous stenting may be performed with braided stainless stents across the inguinal crease with no stent fractures or narrowing due to external compression or hip joint movement. Patency is not associated with the sub-inguinal site of stents, but related to the etiology of the obstruction with secondary patency depending on presence and severity of post-thrombotic obstructions. Iliac vein stents can be safely extended across the groin crease to ensure adequate inflow without fear of compression, fracture or thrombosis related to hip flexion.

10:00 am 26 Reinterventions After Venous Stenting For Chronic Venous Disease

S. Raju¹, P. Tackett², P. Neglén² - ¹University of Mississippi Medical Center, Jackson, MS, ²River Oaks Hospital, Flowood, MS

Background: Percutaneous ilio-femoral venous stenting has been shown to be effective, safe, and durable in both primary and postthrombotic disease. The objective of this study is to analyze those patients that required re-intervention.

Methods: Femoro-ilio-caval stenting was performed in 1060 limbs over a 9 year period (primary/ postthrombotic limb ratio = 555/505). Patients were followed clinically. Stent patency and rate of in-stent restenosis was assessed by venography and ultrasound scanning.

Results: Reinterventions were required in 171 limbs (16%) (11% primary and 16% postthrombotic limbs (p=0.06). Median age, sex ratio, limb side and time of reintervention after the initial procedure (13 months) were similar regardless of etiology of obstruction. Indication for reintervention: swelling 42%, pain 12% or combination 21%, stasis dermatitis/ulcer 12% or stent occlusion 13%; residual or recurrent swelling was more common for primary limbs, pain more common in postthrombotic limbs. Stent occlusion only occurred in postthrombotic limbs; seven had successful thrombolysis and two thrombectomy. Proximal stent extension into the IVC was required in 16 primary limbs due to distal migration of the original stent, and in 12 postthrombotic limbs due to initial inadequate coverage of a postthrombotic IVC segment. Distal extension of the original common iliac vein stent was required in 39 primary limbs to cover previously overlooked non-thrombotic lesion of the external iliac vein in 18 and retroinguinally in 19 limbs. In 30 postthrombotic limbs, distal stent extension was required into external iliac vein (5 limbs) and common femoral vein (25 limbs) to cover postthrombotic strictures. Balloon dilatation of the in-stent restenosis was performed in 57% of limbs; in combination with stent extensions in 32% of limbs. Two types of in-stent restenosis were found, a soft lesion due to layered thrombus and a hard, more fibrous lesion.

Conclusions: Venous stenting for chronic venous disease is remarkably trouble free, with only a small fraction requiring reinterventions. Reintervention were performed to correct overlooked or new defects in inflow, outflow, and/or the stent. Soft in-stent lesions typically occur when inflow or outflow is compromised. The harder lesion occurs independently and is amenable to successful dilation with high pressure balloons. An aggressive stance in identifying and correcting pathology during the initial procedure may reduce the need for reinterventions and improve outcome.

10:20 am **27 Withdrawn**

10:20 am	Venous Stent Registry Update BK Lal, MD
10:25 am	AVF Update – Where the Forum Is Going Mark H. Meissner, MD
10:30 am	Founders Award (TBA) Presented By: Mark H. Meissner, MD
10:35 am	2007 Award Update Introduced By: Mark H. Meissner, MD
	2007 BSN Jobst Winner – Report Danny Vo, MD, Mayo Clinic
10:45 am	2007 Servier Traveling Fellowship Winners – Report Brian Knipp, MD, University of Michigan
10:55 am	2007 Sigvaris Fellowship – Announcement David Gillespie, MD for Reagan Quan, MD, Walter Reed Army Medical Center
11:00 am	PRESIDENTIAL ADDRESS Mark H. Meissner, MD Introduction By: Joann M. Lohr, MD
12:00 pm	MEMBER BUSINESS LUNCH
Free Afternoon	Golf & Tennis

SATURDAY, FEBRUARY 23, 2008

7:00 am	Continental Breakfast – Visit Exhibits
8:00 am	SCIENTIFIC SESSION VI: VENOUS THROMBOEMBOLISM Moderators: Joseph Caprini, MD & Peter Henke, MD
	Educational Objectives:
	1. Upon completion of this session attendees will be able to:
	 Understand the utilization of Duplex ultrasound in defining temporal changes of venous thrombi.
	3.Understand the potential usefulness of combining d-dimer and lower extremity Duplex ultrasound tests in predicting outcome for venous thromboembolism in high risk patients undergoing surgery
	 Understand the complex and controversial management of pregnant females that had previous illo-caval stenting.
	5. Understand the benefits and limitations of diagnostic modalities in patients with suspected venous thromboembolism
	6.Understand the indications, strategy, technical aspects, pharmacologic drugs, mechanical application, and outcomes in treating patients with iliofemoral deep venous thrombosis with pharmacomechanical thrombolysis.
8:00 am	28 Time-Course Analysis of Venous Thrombus With Ultrasonographic Tissue Elasticity Imaging - Preliminary Findings
	K. Uno ^{1,5} , A. Tonomura ² , T. Osaka ² , T. Mitake ² , M. Suda ³ , M. Yamakawa ³ , Y. Isaka ⁴ , S. Homma ⁵ , T. Shiina ³ , K. Aonuma ⁵ - ¹ Namegata District General Hospital, Namegata, Japan, ² Ultrasound Systems Division, Hitachi Medical Corporation, Kashiwa, Japan, ³ Graduate School of System and Information Engineering, University of Tsukuba, Tsukuba, Japan, ⁴ Medical Branch, Academic Service Office for Comprehensive Human Sciences, University of Tsukuba, Tsukuba, Japan, ⁵ Cardiovascular Division, Institute of Clinical Medicine, Graduate School of Comprehensive Human Science, University of Tsukuba, Japan

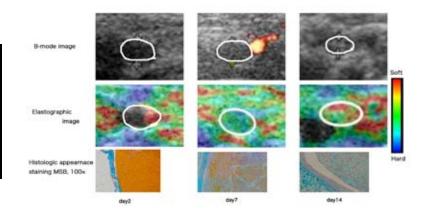
Background: With increasing age venous clots undergo an organization process during which they become adherent to the vessel wall. Therefore, one important factor influencing the decision whether or not to perform thrombolysis or thrombectomy for venous thrombosis is the age of the thrombus. Tissue elasticity imaging is a technology for imaging tissue hardness information using ultrasound. The aimof this study was to examine the diagnostic ability and an appropriate assessment procedure of this method for venous thrombus.

Methods: Conventional Ultrasonography (US) and US-elastography were performed in 25 patients diagnosed venous thrombosis in the lower extremities. All images were obtained with Ultrasound Scanner EUB-8500 (Hitachi Medical Corporation) and analyzed by an external personal

computer. First of all, we examined the elasticity image depending on the different ways of compression and drew a certain assessment procedure for evaluating venous thrombus. Secondly, we observed the venous thrombus with B-mode image, color Doppler image and elasticity image. We estimated the age of thrombus by these conventional ultrasonographic appearances and divided into three phases as follows: acute, sub-acute and chronic. Moreover, we compared the elasticity image with histological evaluation using a rat-based model by ligation of infrarenal Vena cava.

Results: 25 patients, 30 lesions (6 femoral vein; 1 popliteal vein; 22 calf vein; 1 superficial vein) were assessed. We could obtain the stable image by vibrating after initial compression to the area of the thrombus, and setting the region of interest (ROI) including muscle and excluding bone and artery. Mostly, acute phase of thrombi were represented with soft elasticity images, and chronic phase of venous thrombi were represented with hard elasticity images. Meanwhile, acute phase of thrombi with a small amount of venous flow were difficult to differentiate fresh thrombi and venous flow. By histological examination using (figure), US-elastography demonstrated a harder image correlated with the increasing of fibroblast and collagen production in the clots, and recanalisation site showed a soft image.

Conclusions: Venous thrombi were imaged clearly with appropriate compression by US-elastography. We will continue to improve the elasticity image for the application of venous thrombus.



8:20 am 29 Do Preoperative D-Dimer Testing and Venous Duplex Scanning of the Lower Extremities Alter the Outcome In Patients At High Risk For Postoperative Venous Thromboembolism?

T. Yamaki¹, M. Nozaki¹, H. Sakurai¹, M. Takeuchi², K. Soejima³, T. Kono¹ - ¹Tokyo Women's Medical University, Tokyo, Japan, ²Nihon University, Tokyo, Japan, ³Tokyo Metropolitan Hiroo General Hospital, Tokyo, Japan

Background: To assess whether preoperative D-dimer testing and venous duplex scanning (VDS) of the bilateral lower extremities are necessary to identify patients at risk of postoperative venous thromboembolism (VTE).

Methods: One hundred and twenty-seven consecutive referral patients at high or highest risk for postoperative VTE according to the American College of Chest Physicians guidelines were evaluated using pretest clinical probability (PTP) score and D-dimer testing before VDS. After calculating PTP score, patients were divided into low risk (≤ 0 points), moderate risk (1 to 2 points), and high risk (≥ 3 points) PTP. After preoperative VDS was done, all patients received unfractionated heparin or low molecular weight heparin for postoperative VTE prophylaxis. Postoperative surveillance was also performed after operation.

Results: Of 127 referrals, orthopedic patients were the most predominant (42 patients, 33%), followed by gynecologic (30 patients, 24%) and general surgical (17 patients, 13%) patients. Sixty-eight (54%) patients were classified as low, 39 (31%) as moderate, and 20 (15%) as high PTP. Preoperative VDS identified 42 (33%) patients with DVT. Of these, 15 (12%) patients had proximal DVT and remaining 27 (21%) had distal DVT. The prevalence of DVT increased as the risk increased (7%, 56% and 75%, respectively). In the low risk PTP, D-dimer testing provided 80% sensitivity and 97% negative predictive value (NPV) in the diagnosis of DVT. Similarly, in the moderate PTP, the D-dimer testing showed 100% sensitivity and 100% NPV. In the high risk group, D-dimer testing achieved 100% sensitivity and 100% NPV in the diagnosis of DVT. Postoperative VTEs were found in 4 (3%) patients. Three orthopedic patients with low PTP who had preoperative normal D-dimer with no DVT developed symptomatic calf DVTs. One gynecologic patient with moderate PTP who had initial elevated D-dimer with no DVT developed pulmonary embolism. No propagation of DVT or new thrombus formation was found in patients who had preoperative DVT after the operation.

Conclusions: A combination of D-dimer testing and PTP may be effective in detecting patients who require preoperative VDS. However, postoperative VTEs are found predominantly in patients with low PTP, normal D-dimer and no preoperative DVT. Presence of preoperative DVT identified by VDS is not predictive of postoperative propagation of DVT or new VTE formation. Preoperative D-dimer testing and VDS of the bilateral lower extremities do not identify patients at risk of developing clinically important thromboembolic events.

8:40 am 30 Management of Pregnancy In Women With Previous Ilio-Caval Stenting

O. Hartung - CHU Nord, Marseille, France

Background: Ilio-caval stenting represent nowadays the first line treatment for disabling obstructive ilio-caval lesions. Most of the patients are young women which can procreate. We herein report our experience of pregnancy in women who had history of ilio-caval stenting.

Methods: From November 1995 to July 2007, 112 patients had ilio-caval stenting for obstructive venous disease in our department. Of these, 61 women were not menopaused. Seven pregnancy occurred in five patients (mean age 25 years) after stent deployment (one patient had 3 pregnancy). They had stenting for May-Thurner disease in 3 cases and during venous thrombectomy in 2 cases. All stents were self-expanding metallic stents located on the left common iliac vein. All of them received preventive treatment with low molecular weight heparin (LMWH) from the third month of pregnancy to one month after delivery and had to wear elastic stockings. Patients also had to sleep on their right side if possible. They were followed during the pregnancy by dupplex-scan at 3, 6, 8 months and then one month after delivery.

Results: One patient had unrelated spontaneous abortion after 2 month of pregnancy. No deep venous thrombosis nor symptomatic pulmonary embolism occurred during pregnancy, delivery and post-partum. Four patients needed cesarean delivery and none had hemorrhagic complication. None of the patients had adverse effects of the treatment. Dupplex-scan showed compression of the stent(s) at 8 months in 3 cases with inflow obstruction in 2 cases.Post partum dupplex-scan showed in all cases that stents were patent with no remaining stenosis. No stent had structural damage.

Conclusions: Ilio-caval stent(s) compression can occur during pregnancy but do not lead to structural damage of self-expanding stents. Despite this no deep venous thrombosis occurred with preventive LMWH treatment.

9:00 am

31 The Evaluation of Diagnostic Procedures of Venous Thromboembolism (VTE) In Patients With Suspected VTE

J. Lee, B. Zierler - University of Washington, Seattle, WA

Background: Evidences-based guidelines for VTE diagnosis recommend that pre-test probability (PTP) of VTE and D-dimer before any objective diagnostic tests be performed as the initial screening test particularly in emergency room or out-patient clinics. The purpose of this study was to evaluate VTE diagnostic procedures in patients with suspected VTE prior to the implementation of the VTE Safety Toolkit.

Methods: The study was a part of a larger pre/post study to implement a VTE Safety Toolkit consisting of clinical algorithms for the prevention, diagnosis, and management of VTE. A vascular laboratory logbook and medical records of adult patients who underwent lower extremity venous duplex scans (VDS) during the 6-month pre-intervention period were retrospectively reviewed in order to identify the utilization patterns of VTE diagnostic tests including D-dimer, VDS for deep vein thrombosis (DVT), and ventilation and perfusion (VQ) scan or computerized tomographic (CT) angiography for pulmonary embolism (PE). We developed DVT/PE diagnostic algorithms as a component of the VTE Safety Toolkit. We made a determination using the diagnostic algorithm of whether the number and order of objective tests were appropriate given the VTE risk score and patient history.

Results: Approximately 972 lower extremity venous duplex scans in 818 patients with suspected VTE were performed in 6 months of the pre-intervention period at an academic medical center. Among the 818 patients, 112 patients (13.7%) were diagnosed with DVT by VDS. A quarter of patients with VDS were asymptomatic and 16 % (32/203) of these patients were diagnosed with DVT. Approximately 27% (30/112) of patients with acute DVT had serial VDS and propagation was identified in 9 patients with follow-up VDS (30%, 9/30). We categorized VTE diagnostic strategies into four; 1) duplex only, 2) D-dimer and duplex, 3) D-dimer/duplex/lung scanning by CT or VQ scans, and 4) duplex/CT or VQ scan. The rates of diagnosis of DVT/PE were higher in patients in the strategy included D-dimer ± Duplex ± CT/VQ compared to those in other strategies (please see the table below).

Conclusions: There was inappropriate utilization of VTE diagnostic tests. The VTE Safety Toolkit includes a tool requires referring providers to rate the PTP before obtaining an objective study for VTE diagnosis. The VTE Safety Toolkit diagnostic algorithms should increase the appropriateness of the diagnostic studies ordered.

Outcomes by VTE diagnostic strategies		1. Duplex only	2. D-dimer ± Duplex	4. Duplex ± CT/VQ	
		535/818 (65.4%)	121/818 (14.8%)	102/818 (12.5%)	
DVT diagnosis (1	12/818, 13.7%)	62/536 (11.6%)	11/121 (9.1%)	24/102 (23.5%)	
PE diagnosis					
(58/818, 7%)		0	0	34/102 (33.4%)	
Indications for Du	Frequency (%)				
1. Rule out DVT (v pain)	568/818 (69.4%)				
2. Look for source shortness of brea	184/818 (22.5%)				
3. Surveillance (w prolong bed rest	100/818 (12.2%)				
4. Asymptomatic	203/818 (24.8%)				

9:20 am

32 The Quantitative Benefit of Isolated, Segmental, Pharmacomechanical Thrombolysis For Iliofemoral DVT

J. Martinez, A. J. Comerota, S. Kazanjian, R. DiSalle, D. M. Sepanski, Z. I. Assi - The Toledo Hospital, Toledo, OH

Background: It is becoming increasingly recognized that early thrombus removal in patients with iliofemoral DVT reduces postthrombotic morbidity. Preserving valve function and relieving venous obstruction prevents deterioration of quality of life and loss of economic potential. The preferred method for treating iliofemoral DVT is catheter-directed thrombolysis (CDT). Recently, isolated segmental pharmacomechanical thrombolysis (ISPMT) has emerged as a treatment option for patients with extensive DVT. This technique isolates the venous segment being treated between two occluding balloons and delivers the plasminogen activator into the thrombus. The catheter assumes a spiral configuration and rotates at 1500 RPM, macerating the thrombus. Subsequently, the liquefied thrombus and remaining lytic agent are aspirated. The purpose of our study is to determine whether there are advantages to ISPMT and, if so, to quantify those advantages relative to CDT.

Methods: Forty-three patients treated with CDT at our institution between May 2003 and June 2007 were reviewed. Patients were divided into two groups, those treated with ISPMT (Trellis® 8 catheter, Bacchus Vascular, Santa Clara, CA) or those using CDT alone. Data obtained included demographics, extent of thrombus, procedural details, periprocedural evaluation, and thrombus resolution. CDT was performed by placing a multi-hole catheter directly into the thrombus; infusion was administered for varying periods of time, and repeat phlebography was used to determine treatment time. Amount of lysis was determined by comparison of pre- and post-procedure phlebographic images.

Results: Catheter-directed thrombolysis alone was used in 21 patients, and 22 had ISPMT incorporated as part of the strategy of thrombus removal. Treatment time (52.2 vs. 22.9 hrs; P=0.0001) and dose of rt-PA (55.3 vs. 32.5 mg; P=0.007) were decreased by the use of ISPMT. The use of ISPMT improved overall lytic success (84.3% vs. 92.3%; P=0.029) and more patients had complete thrombus resolution (Table). There was no difference in the use of adjunctive therapy such as venoplasty with or without stenting. Complication rates were similar and there was no major morbidity or mortality. Hospital length of stay and ICU length of stay were similar between the two groups.

Conclusions: The use of ISPMT offers more effective thrombus removal in less time and with a reduced dose of thrombolytic agent. However, decreased treatment time did not translate into decreased hospital or ICU stay. Longer-term follow-up is required to determine whether improved thrombus resolution translates to better functional outcome and reduced postthrombotic morbidity.

Percent of lysis by group					
% Lysis CDT ISPMT					
Complete (≥95)	5/21 (23%)	15/22 (68%)			
Significant (75-94)	12/21 (57%)	5/22 (23%)			
Moderate (50-74)	4/21 (19%)	2/22 (9%)			
Minimal (<50)	0/21	0/22			

9:40 am

Coffee Break / Visit Exhibits

10	:1	0	am

SCIENTIFIC SESSION VII: CHRONIC VENOUS DISFASE II

Moderators: Joann Lohr, MD & Fedor Lurie, MD

Educational Objectives:

- 1. Upon completion of this session attendees will understand:
- 2. The implications of chronic venous disease and venous hypertension on the adverse effect on arterial hemodynamics.
- 3. The risk factors implicated in patients with varicose veins that will progress to venous ulceration.
- 4. The risk factors associated with recalcitrant venous ulcer treated with compression.
- 5. The technical application of a neovalve for deep venous insufficiency, comparing two different experiences.

10:10 am

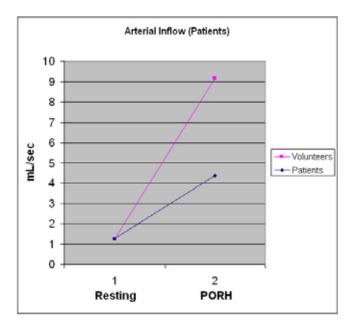
33 Lower Extremity Arterial Inflow Is Adversely Affected In Patients With Postthrombotic Venous Disease D. Paolini, L. Jones, A. J. Comerota - The Toledo Hospital, Toledo, OH

Background: Chronic venous disease of the lower extremity is due to venous hypertension resulting from reflux and/or obstruction. Studies in patients with postthrombotic venous disease with ambulatory leg discomfort have focused primarily on venous hemodynamics of the lower extremity. The impact of postthrombotic venous disease on arterial perfusion has not been evaluated. The purpose of this study is to evaluate whether arterial perfusion at rest and following stress induced by postocclusive reactive hyperemia (PORH) is adversely affected by postthrombotic venous disease.

Methods: Lower leg arterial perfusion was measured in 16 limbs with C3-C6 postthrombotic venous disease (study group) and compared to 22 disease-free limbs (control group). Neither group had any history, signs, or symptoms of arterial occlusive disease, and all had a normal ankle-brachial index at rest. Arterial perfusion was measured, using venous occlusive plethysmography, at rest and following 3 minutes of complete leg ischemia induced by a thigh cuff inflated to 20 mmHg above systolic pressure.

Results: Resting arterial inflow was similar in control and study groups: 1.26 mL/sec and 1.28 mL/sec (P=NS), respectively. PORH stimulated a 7.3-fold increase in arterial inflow to 9.17 mL/sec in the control group, whereas the study group increased only 3.4-fold to 4.38 mL/sec (P=0.0045) (Figure).

Conclusions: Arterial inflow is significantly diminished after PORH in patients with postthrombotic venous disease. These data suggest that the etiology of ambulatory leg pain in patients with postthrombotic venous disease is at least in part related to a reduction of arterial inflow during exercise.



10:30 am

34

Which Patients With Varicose Veins Are At Increased Risk of Chronic Leg Ulceration?

L. Robertson¹, A. Lee², K. Gallagher³, S. Carmichael³, C. Evans⁴, B. McKinstry¹, S. Fraser³, P. Allan¹, C. Ruckley¹, F. Fowkes¹ - ¹University of Edinburgh, Edinburgh, United Kingdom, ²University of Aberdeen, Aberdeen, United Kingdom, ³Lothian University Hospitals NHS Trust, Edinburgh, United Kingdom, ⁴NHS Lothian, Edinburgh, United Kingdom

Background: When prioritising clinical management there is a need to identify which patients with varicose veins will go on to develop chronic leg ulcer. This is the first study to report, in patients with varicose veins, the characteristics of venous disease and lifestyle factors related to an increased risk of ulceration.

Methods: A case control study compared 120 cases with varicose veins and an open or healed leg ulcer, with 120 controls with varicose veins but no leg ulcer. Assessment included: clinical classification of venous disease, duplex scanning, quantitative digital photoplethysmography, body mass index (BMI), and questionnaire on social class, smoking, current and previous leisure exercise, mobility at work and previous history of venous disease.

Results: Cases had a higher mean age than controls (61.1 years (13.4) versus 59.9 years (11.7), p=0.01). Severity of venous disease was linked to increased risk of ulceration. Other significant risk factors included history of deep vein thrombosis or pulmonary embolism (OR 4.1, 95% CI 1.8-9.7), increased BMI (OR 1.1, 95% CI 1.0-1.1) and smoking (OR 2.0, 95% CI 1.2-3.8), and remained so after adjusting for age and sex. Cases had reduced calf muscle pump power and limited range of ankle movement (not wholly due to the effects of an active ulcer) (both p<0.05). Reflux in the profunda, femoral and popliteal veins were also significantly associated with increased risk of ulceration.

Risk factors that were significant on univariate analyses were entered into a stepwise logistic regression model to determine which factors were independently associated with increased risk of ulceration.

Risk factor	Multi-adjusted odds ratio (95% CI)
BMI	1.08 (1.01-1.15)
Venous pump power	0.96 (0.92-0.99)
Dorsiflexion	0.88 (0.81-0.97)
Corona phlebectatica	4.52 (1.81-11.3)
Eczema	2.82 (1.12-7.07)
Lipodermatosclerosis	8.90 (1.44-54.8)
Reflux in popliteal vein	2.82 (1.03-7.75)

Conclusions: The results of this study confirm that, in patients with varicose veins, those with skin changes of chronic venous insufficiency and deep vein incompetence, are at greatly increased risk of ulceration. However, the risks may also be increased in those who smoke, are obese, and have conditions restricting ankle movement. A prognostic scoring system incorporating all these factors is being developed to assist clinical decision making.

10:50 am

35 Risk Factors Related To the Failure of Venous Leg Ulcers To Heal With Compression Treatment

D. J. Milic, S. S. Zivic, D. C. Bogdanovic, V. D. Milojkovic, M.A. Pejic, V. M. Popovic - Clinic for Vascular Surgery, Clinical Centre Nis, Nis, Serbia

Background: Compression therapy is the most widely used treatment for venous leg ulcers and it was used in different forms for more than 400 years. Published healing rates of venous ulcers obtained with compression therapy vary widely from 40-90%. According to numerous studies it has been suggested that the application of external pressure to the calf muscle raises the interstitial pressure resulting in improved venous return and reduction in the venous hypertension. Several risk factors have been identified to be correlated with the failure of venous leg ulcers to heal with compression therapy (longer ulcer duration; large surface area; fibrinous deposition present on >50% of the wound surface and an ankle brachial pressure index of <0.85).

Methods: An open prospective, single-center study was performed in order to determine possible risk factors associated with the failure of venous ulcers to heal when treated with multi-layer high compression bandaging system for 52 weeks. One hundred and eighty nine patients (101 women, 88 men; mean age 61 years) with venous leg ulcers (ulcer surface >5cm2; duration >3 months) were included in the study. The study excluded patients with arterial disease (ABPI<0.8), heart insufficiency with EF<35, pregnancy, cancer disease, rheumatoid arthritis and diabetes. Based on clinical opinion and available literature, the following were considered as potential risk factors: sex, age (years), ulceration surface (cm2), time since ulcer onset (months), previous operations (stripping, SEPS), history of deep vein thrombosis, body mass index (BMI), microbiological status of the wound, reduction in calf circumference, walking distance during the day, calf: ankle circumference ratio <1.3, fixed ankle joint, ulcer recurrence, number of wounds, history of wound debridement, >50% of wound covered with fibrin, lipodermatosclerosis, depth of the wound >1.5 cm.

Results: Within 52 weeks of limb-compression therapy, 24 (12.7%) venous ulcers had failed to heal. A small ulceration surface (< 20 cm2), the duration of the venous ulcer <12 months, a decrease in calf circumference of more than 2 cm during the first 50 days of treatment and emergence of new skin islets on the wound surface were favorable prognostic factors for ulcer healing. A large BMI (>35 kg/m2), short walking distance during the day (<200m), a history of wound debridement and ulcers with deepest presentation (>2cm) were indicators of slow healing. Calf: ankle circumference ratio <1.3, fixed ankle joint and reduced ankle range of motion (<20 degrees) were the only independent parameters associated with non-healing (P<0.01).

Conclusions: The results obtained in this study suggest that non healing venous ulcers are related to the impairment of the calf muscle pump.

11:00 am 36 Neovalve Construction In Deep Venous Incompetence: Comparison Between Two Subsequent Case Series and Related Technical Details

M. Lugli, S. Guerzoni, O. Maleti - Hesperia Hospital, Modena, Italy

Background: Deep venous incompetence, due to postthrombotic syndrome and primary valvular defects, is a main cause of severe chronic venous disorders, often resistant to conservative therapy. In selected cases deep venous surgery should be considered and when more consolidated techniques are not suitable, neovalve construction by parietal dissection can be performed. This technique was applied in two subsequent case series, which differ in surgical technical details after identifying one probable cause of valve failure in the first group.

Methods: From December 2000 to June 2007 we performed 40 neovalve construction operations in 36 patients (19 males, 17 females, median age 57, range 29 - 82) affected by deep venous insufficiency. 32 patients were affected by post thrombotic syndrome and 4 by valve agenesia. The 32 patients with post-thrombotic syndrome were selected from among 76 with resistant ulcers classified C 6,S E S A S,D,P P R,RO and the 4 patients with valve agenesia were selected within 28 affected by resistant ulcers classified as C 6,S E P A S,D,P P R. The patients were subdivided into 2 groups: the first included 19 operations performed in the period from December 2000 and December 2004, presenting a mean follow up of 57 months (range 31-78) and the second group included 21 patients from January 2005 to June 2007 with a mean follow up of 11 months (range 2-29). In the second group a surgical variation was applied in order to prevent flap collapse and improve neovalve competence. The principal technical variation consists in a modification of the flap principally by performing a fixation of it in the semi-open position. Postoperative venography was performed in all cases.

Results: In the first series ulcer healing was observed in 16 cases out of 19 (84%). Recurrent ulcers were observed in one case after three years. The valvular continence was found in 13 out of 19 (68%). Regarding the second series, continence was obtained in all cases. The ulcer didn't heal in one case (4.8%) and recurred in two cases (9.5%). Postoperative deep venous thrombosis was observed in 3 patients in the first series and was not detected in the second. Mortality rate was 0 and pulmonary embolism was not detected in both groups.

Conclusions: The modified technique applied to the second group seemed to improve the valve continence significantly. However a longer period of follow up for this latter group is required to validate this technical enhancement.

11:30 am

D. EUGENE STRANDNESS MEMORIAL LECTURE

Introduced by: Mark H. Meissner, MD

12:30 – 1:30 pm	INDUSTRY SPONSORED LUNCHEON
	ClosureFAST™ Clinical Trials Update
	By: VNUS Medical Technologies
	1) Prospective, multicenter 1 year follow-up
	2) Randomized trial comparing ClosureFAST to endovenous laser
	3) Lessons learned 1 year post launch
1:30 pm	ASK THE EXPERTS: PELVIC CONGESTION SYNDROME Moderator: Chieh Min Fan, MD
	Educational Objectives:
	 Review anatomy and techniques for imaging the venous structures of the pelvis and retroperitoneum.
	2. Recognize the clinical manifestations of pelvic venous congestion
	3. Understand endovascular and surgical treatment approaches for pelvic venous congestion syndromes
	4.Review classification, clinical patterns, and treatment approach for pelvic vascular anomalies
	Anatomy and Imaging of the Venous System of the Pelvis and Retroperitoneum Chieh Min Fan, MD, Brigham and Women's Hospital, Boston, MA
	Pelvic Congestion Syndrome: Diagnosis and
	Management Anthony C. Venbrux, MD, George Washington U. Hospital, Washington, DC
	Nutcracker Syndromes: Endovascular and Surgical Management

Matthew Menard, MD, Brigham and Women's Hospital, Boston, MA

Vascular Anomalies: An Uncommon Cause of Pelvic Venous Congestion

Patricia Burrows, MD, St. Lukes – Roosevelt Hospital Center, New York, NY

2:30 pm Coffee Break / Visit Exhibits

HOW TO SESSION

Recanalization and Re-Endovenous Ablation; Mapping Out My Veins Tips and Tricks; Sclero the Do's and Don'ts

Moderator: Julianne Stoughton, MD

Educational Objectives:

- Attendees will become familiar with many of the common, and some of the unusual (but important) patterns of venous anatomy
- 2.There will be a discussion involving the approach to incompetent perforating veins: reviewing the treatment options, the technical aspects of each treatment, as well as a discussion of which veins are best treated with which technology
- 3.Recannalization, neovascularization and recurrent veins after venous intervention and will be discussed. The approaches will be illustrated with case presentations
- 4.Difficult management cases will be presented including: the hypercoaguable patient, the obese patient, patients with anomalous anatomy, etc.

Endovenous Heat Induced Thrombosis: When To, How To and What To Look For Lowell Kabnick, MD

Treatment of Incompetent Perforators Steve Elias, MD

Treatment of Neovascularization and Recannalized Veins

Ronald Bush, MD

MODERATED POSTER SESSION

Moderator: Michael Dalsing, MD Frank Padberg, MD Bo Eklof, MD

Educational Objectives: The participants in the poster session will gain a wide range of knowledge expansion including chronic venous disorder, saphenous vein treatment, understanding risk factors and evaluation methods.

P-1 Microparticles: A Natural History Time Course Analysis In A Model of Murine Venous Thrombosis

A. E. Hawley, D. M. Farris, N. E. Ballard, A. P. McDonald, S. K. Wrobleski, P. K. Henke, D. D. Myers, T. W. Wakefield - University of Michigan, Ann Arbor, MI

Background: Deep venous thrombosis remains a significant health care problem. Recent studies suggest that procoagulant microparticle (MP) formation plays an integral part in the inflammatory component of venous thrombosis, leading to thrombus amplification. Low levels of MPs are found circulating in a normal physiological state and have been shown to increase with disease. The purpose of this study was to document a detailed time-course analysis of MPs after the initiation of venous thrombosis, and to assess associated tissue factor (TF) expression.

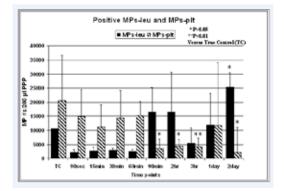
Methods: Inferior vena cava (IVC) ligation was performed on C57BL/6 mice (n65) utilizing an established model of IVC occlusion. Animals were sacrificed, post ligation, at the following time points (90sec, 15min, 30min, 60min, 90min, 2hour, 3hour, 1day, 2day) to assess MP production and thrombus mass. Approximately 500µL of anti-coagulated blood was used to obtain platelet poor plasma (PPP). The PPP was centrifuged and the pelleted MPs measured by flow cytometry (Becton Dickinson FACS Calibur with Cell Quest software). MPs were stained which identified them as originating from either leukocytes (MP-leu) or platelets (MP-plt). An additional MP pellet, pooled from 2day animals (n40), was evaluated at varying concentrations for TF procoagulant activity by chromogenic assay.

Results: MPs-leu and MPs-plt were significantly decreased vs. True Controls (TC) at 3hour. On Day 2, MPs-leu showed a significant increase while MPs-plt remained decreased significantly vs. TC (Figure 1). Thrombus mass correlated positively with MPs-leu, and negatively with MPs-plt (Table 1). TF bearing MPs showed a direct relationship to MP concentrations (R=0.99). Animals with IVC occlusion, but no clot, averaged 40% less TF (Table 2).

Conclusions: MPs tend to decrease after thrombosis. However, those from leukocyte origin increase significantly by day 2, while those from platelet origin remain depressed, likely from consumption into the thrombus. Thrombosis correlates positively with MPs from leukocyte origin. Additionally, MPs demonstrate a highly significant positive correlation with TF activity. This study suggests that knowing the time course and origin of MPs is important to understanding their biology and their potential use as a diagnostic technique.

Table 1:	
Groups	Average IVC/Thrombus Mass (grams/cm)
True Controls	.0045
90sec	.0046
15min	.0061
30min	.0053
60min	.0041
90min	.0063
2hour	.0077
3hour	.0154
1day	.0179
2day	.0256
R=0.65(Mass vs. Mps-leu)	R=-0.59(Mass vs. Mps-plt)

Table 2:	
Tissue Factor Activity of Microparticles	
# of MPs	pM TF avg.
10000	13.7
25000	40.5
50000	75.9
80000	124.3
160000	205.2
No clot 80000	64.6
No clot 160000	144



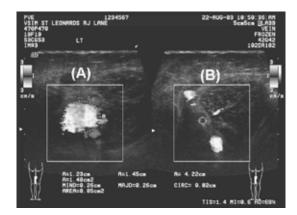
P-2 Popliteal Vein Compression Syndrome: Obesity, Venous Disease and the Popliteal Connection R. J. Lane¹, M. L. Cuzilla² - ¹Royal North Shore Hospital, Sydney, Australia ²Vascular Surgery Inventigations and Managment, Sydney, Australia

Background: Obesity and venous disease are commonly encountered together. The aetiological relationship however has not been clear. Popliteal venous compression (PVC) has been encountered both on ultrasound and venographically. In this study, patients with symptoms and/or signs of chronic venous hypertension with PVC were investigated and the relationship to obesity defined. Popliteal Vein Compression Syndrome (PVCS) is defined herewith in its pathological state.

Methods: Colour duplex ultrasound (CDU) was used to measure the maximum internal diameter (ID) of the popliteal vein (POPV) with the knee locked and unlocked. A total of 89 patients were included in the study of which 49 limbs were classified as having PVC as determined by colour duplex ultrasound. A positive finding was defined as a greater than 90% reduction in the maximum internal diameter (ID) of the POPV with knee locking. Forty consecutive limbs with venous disease with no evidence PVC were used as controls. The Body Mass Index (BMI) of each group was calculated and the clinical symptoms and signs were documented. After failed conservative treatment, 30 of the 49 underwent open popliteal decompression.

Results: Patients with PVC were found to have a BMI of 36.0 ű 6.2 compared with the controls of 25.3 ű 3.0. The POPV ID in the PVC group before and after knee locking changed from 11.7 ű 5.0 mm to 1.0 ű 2.1mm respectively. Post-operatively, the POPV ID before and after knee-locking changed from 10.2 ű 2.2 mm to 9.0 ű 1.5 mm respectively. At 16.2 ű 12.1 months follow-up, all the major clinical parameters including pain, oedema, pigmentation and lipodermatosclerosis plus the total clinical score were all improved at a statistically significant level.

Conclusions: There appears to be a relationship between obesity, chronic venous disease and PVC. Popliteal vein compression syndrome may clarify the previously unexplained venous presentations, including worsening following ablative venous procedures, and perhaps the known association with deep vein thrombosis and obesity. Surgical decompression provides gratifying results in patients unresponsive to conservative treatment.





P-3 Ultrasonic Venous Valve Imaging - A Prerequisite For Exostent Repair

R. J. Lane¹, M. N. Phillips², M. L. Cuzilla³ - ¹Royal North Shore Hospital, Sydney, Australia; ²AllVascular Pty Ltd, Sydney, Australia; ³Vascular Surgery Inventigations and Managment, Sydney, Australia

Background: Lower limb venous disease remains a significant problem in our community today. The condition has been treated mainly with ablative procedures such as stripping and or sclerotherapy. The aim of this study was to define the ultrasonic features of repairable venous valves by External Valvular Stenting (EVS). In addition, to access the ability to predict success of EVS determined intra-operatively and at three-months post-operatively.

Methods: Valves considered for EVS were assessed with Brightness-Mode (B-Mode), Spectral Pulsed Doppler (PD), Colour Doppler Imaging (CDI) and Brightness-Flow (B-Flow).

The ultrasonic features of the great saphenous vein (GSV), terminal valve (TV) and sub-terminal valves (STV) were considered. Inclusion criteria were; valvular ring dilation <12mm in diameter, (GSV) internal diameter (ID) <12mm along the entire length of the trunk, symmetry of the valve sinuses, positive identification of two valve cusps, and symmetrical reflux flow patterns through the incompetent valve. There were 69 limbs included in the study. All repaired TV's were tested intra-operatively for competence after application of the EVS. If there was evidence of residual reflux, the STV was also repaired. The operated limbs were assessed clinically 3 months after the procedure at which time ultrasound was also used to test the repaired valves.

Results: Of the 69 TV's that were examined pre-operatively, a total of 50 were considered repairable by ultrasonic features (72%). At operation, 44 of these valves were successfully repaired (88%). In the 6 limbs which had residual TV reflux, the STV was repaired. All 6 had competence in the GSV trunk following the STV EVS. Of the 19 TV's that were considered by ultrasonic features to be unrepairable, 18 had gross reflux following EVS with 1 only repaired being successful. All limbs that were successfully repaired at operation were followed up 3 months later, and re-examined with diagnostic ultrasound. Of this group; 3 GSVs had residual reflux at the TV and STV, 1 GSV had major reflux and 1 GSV developed thrombophlebitis. The overall figures for the predictability of successful EVS based on ultrasonic features of the valve were; sensitivity 97.8% (95% CI, 88.2 - 99.6), specificity 75% (95% CI 53.3 - 90.2) and accuracy 90.4%.

Conclusions: In the treatment of varicose veins, a combination of ultrasound modalities accurately predicts EVS outcomes at the TV and STV of the GSV.

P-4 Prevalence and Distribution of Deep Vein Thrombosis In Patients With Symptomatic Pulmonary Embolism

T. Yamaki¹, M. Nozaki¹, H. Sakurai¹, M. Takeuchi², K. Soejima³, T. Kono¹ - ¹Tokyo Women's Medical University, Tokyo, Japan; ²Nihon University, Tokyo, Japan; ³Tokyo Metropolitan Hiroo General Hospotal, Tokyo, Japan

Background: To investigate the prevalence and distribution of deep vein thrombosis (DVT) in patients with symptomatic pulmonary embolism (PE), and to compare characteristics between patients with PE and these without PE.

Methods: A total of 420 consecutive patients with DVT were included. The distribution of DVT was evaluated with duplex scanning, and patients with clinical suspicion of PE were investigated using ventilation/perfusion lung scintigraphy. The patients were then followed for 6 months for investigation of recurrence of venous thromboembolism (VTE) and outcome.

Results: PE was found in 82 (20%) patients. There were no significant differences in mean age, gender, risk factors for VTE, and laterality of leg involvement between patients with DVT and PE and these with DVT alone. On the other hand, the proportion of leg symptoms was statistically higher in patients with DVT alone (p=0.0002). The most common venous segment containing thrombosis was SV in both groups (57% and 50%, respectively, p=0.254). However, there was a significantly higher proportion of distal DVT in patients with DVT and PE (p<0.0001). The significantly higher proportion of DVT was found in gastrocnemius vein in patients with DVT and PE (p=0.018). In contrast, the proportion of common femoral vein thrombosis was found to be significantly higher in patients who had DVT alone (p=0.049). There were similar tendencies in the proportions of distinguished risk factors for patients with DVT alone. There was a significant higher proportion of recurrent VTE and mortality rate in patients with DVT and PE. (p < 0.0001 and p = 0.0042, respectively). In patients with DVT and PE, there was no significant difference in the cumulative proportion of overall survival rate between the patients with proximal DVT and these with distal DVT (p=0.600).

Conclusions: The lower extremity venous duplex scanning demonstrates that distal DVT is more predominant in patients who had PE compared to these with DVT alone. Recurrent VTE is more often found in patients with DVT and PE. Mortality rate is much worse in patients with DVT and PE. However, overall survival rate appears to be similar between PE patient with proximal DVT and these with distal DVT. Although, there are similar tendencies in age, gender, laterality of leg involvement, and risk factors for VTE between patients with DVT and PE and these with DVT alone, these two diseases appear to be distinct with different distribution of DVT and natural history.

P-5 Endovenous Laser Therapy In the Treatment of Short Saphenous Varicose Veins: A Non-Randomised Controlled Trial

A. Mekako, J. Hatfield, S. Gulati, M. Abdul Rahman, P. T. McCollum, I. C. Chetter - Hull Royal Infirmary/University of Hull, Hull, United Kingdom

Background: Endovenous laser therapy (EVLT) is a safe and effective treatment modality for varicose veins on short-to-medium term follow-up. Reports on EVLT have largely focussed on varicosities affecting the greater saphenous vein. This study compares EVLT with surgery in the treatment of short saphenous varicose veins.

Methods: Two non-randomised groups were studied. EVLT (14W continuous): 22 patients, median age 48 (IQR 43-56) years, and surgery (sapheno-popliteal junction ligation +/- strip and avulsions): 18 patients, median age 48 (IQR 30-57) years. Patients were assessed at 1, 6, and 12 weeks post-procedure. Pain scores, venous clinical severity scores (VCSS), abolition of reflux, return to work /normal activities, patient satisfaction, as well as quality of life (QoL) outcomes were analysed.

Results: Baseline parameters were similar. Visual Analogue Scale-rated mean pain scores were significantly lower following EVLT during the first week (1.3 versus 3.2; p=0.003). There were no significant differences in VCSS, return to work /normal activities, abolition of reflux, and patient satisfaction at 12 weeks. The EVLT group had significantly lower mean Aberdeen Varicose Vein Scores at 1 week (15.45 versus 20.29; p=0.01), but no differences at 6 and 12 weeks. There were no differences in SF-36 domain scores, except at 6 weeks (better social functioning following EVLT: 84 versus 74; p=0.04).

Conclusion: This study has demonstrated some benefits of EVLT over surgery in the early post-operative period, although outcomes were largely similar in both groups. This may suggest equivalence of both treatment modalities, but a randomised trial is indicated to elucidate longer-term clinical and QoL outcomes.

P-6 Greater Saphenous Vein Diameter Predicts Venous Reflux

J. Bloom, F. C. Vandy, S. Brown, A. Clay, C. Lane, G. Reynolds, S. LeBaron, C. Nighswander, P. K. Henke, T. W. Wakefield - University of Michigan, Ann Arbor, MI

Background: Current literature has demonstrated a positive correlation with venous diameter and increased valve closure time suggesting reflux. Specifically, this has been shown for perforating veins. However, this association has not been demonstrated for the greater saphenous vein (GSV). This study investigated whether such a relationship exists between valve closure time and vein diameter.

Methods: A random subset of 135 patients (188 limbs) from our endovascular laser therapy (EVLT) registry had undergone a venous reflux study. Patients were placed in reverse Trendelenburg and evaluated for venous incompetence as measured by valve closure time in msec. GSV diameter was measured at the saphenofemoral junction and reflux measurements were obtained 2 centimeters distal to the valve with the Doppler probe in the saggital position and parallel to the vessel wall. Valve closure time in the common femoral and greater saphenous veins was recorded following a Valsalva maneuver. However, valve closure in the popliteal vein was obtained with a distal thigh compression. GSV diameter was compared to venous valve closure time using one-way analysis of variance (ANOVA). Post-hoc analysis was done using a Bonferroni's correction.

Results: Valve closure times were categorized corresponding to the degree of reflux. (Table 1) A valve closure time indicative of moderate reflux was statistically longer than that of mild reflux. (p = .002) There was no statistical difference in valve closure time between normal and mild or moderate and severe. A 2-tailed independent t-test revealed the GSV diameter of a combination moderate and severe group to be significantly larger than a combination mild and normal group. (Table 2) There was no statistical association between GSV diameter and valve closure time in the popliteal or common femoral veins.

Conclusions: Although we did not identify difference between every classification of reflux, there was a statistical difference between mild and moderate classes. A simplified analysis comparing our combined normal and mild groups to our combined moderate and severe groups demonstrates a direct relationship with GSV diameter and reflux time. Our data as well as our clinical observations suggest diameters greater than .795 centimeters are associated with a moderate to severe degree of reflux. These results call into question the clinical significance of a small GSV diameter in the presence of a moderate to severe reflux.

Table 1.						
	N=	% female	% right leg	GSV diameter	Std. Dev	95% CI
Classification						
Normal (0- 1000 msec)	11% (21)	71% (15)	47% (7)	0.6695 cm	0.22128	0.5688- 0.7702
Mild (1001- 2000 msec)	20%(37)	76% (28)	54% (20)	0.6616 cm	0.18986	0.5983- 0.7249
Moderate (2001-3000 msec)	31% (58)	67% (39)	53 (31)	0.8712 cm	0.34574	0.7803- 0.9621
Severe (>3000 msec)	38% (72)	74% (53)	42% (30)	0.8224 cm	0.2403	0.7659- 0.8788

Table 2.					
	N=	GSV diameter	Std. Dev	95% CI	
Classification					
Normal and Mild	58	0.664 cm	0.200	0.612-0.717	
Moderate and Severe	130	0.847 cm	0.293	0.795-0.898	p = 0.000

Posters

P-7 Combined Intermittent Pneumatic Leg Compression and Pharmacological Prophylaxis For Prevention of Venous Thromboembolism In High Risk Patients

S. K. Kakkos¹, J. A. Caprini², G. Geroulakos³, A. N. Nicolaides³, G. P. Stansby⁴, D. J. Reddy¹ - ¹Henry Ford Hospital, Detroit, Ml; ²Evanston Northwestern Healthcare, Evanston, IL; ³Imperial College, London, United Kingdom; ⁴University of Newcastle upon Tyne, Newcastle upon Tyne, United Kingdom

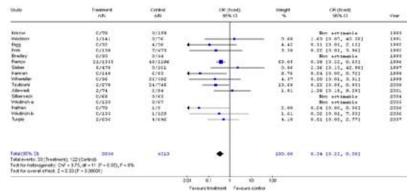
Background: It has been suggested that combined modalities are more effective than single modalities in preventing venous thromboembolism in high risk patients. The aim of the present study was to perform a meta-analysis on the efficacy of intermittent pneumatic leg compression combined with pharmacological prophylaxis versus single modalities (intermittent pneumatic leg compression or pharmacological prophylaxis) in preventing venous thromboembolism in high-risk patients.

Methods: Using MEDLINE search and the reference lists of relevant articles to identify additional trials, studies that used combined intermittent pneumatic leg compression and pharmacological interventions to prevent venous thromboembolism in high-risk patients were identified.

Results: Seventeen studies, six of them randomized controlled trials (RCTs), which enrolled 9,998 patients in total in a variety of specialties, including orthopedic, general and cardiac surgery, were identified. Five RCTs evaluated the role of combined modalities on the incidence of symptomatic PE. These showed a reduction in PE from 2.51% (53/2110) in the control group (single modalities) to 1.03% (24/2335) in the treatment group. Relative risk was 0.41, 95% confidence interval (CI) was 0.25 to 0.65. Results did not demonstrate heterogeneity or publication bias (I2=0%). Five RCTs investigated the role of combined modalities on the incidence of DVT. These showed a reduction in DVT from 4.79% (87/1816) in the control group to 2.29% (43/1881) in the treatment group. RR was 0.43, 95% CI 0.30 to 0.61. Results were consistent with significant heterogeneity (I2=66.2%).

Repeat analysis that included all studies revealed similar results. Sixteen of the included studies evaluated the role of combined modalities on the incidence of symptomatic PE. These showed a reduction in symptomatic PE from 2.83% (122/4313) in the control group to 0.86% (33/3838) in the treatment group (Figure). Odds ratio was 0.34, 95% CI 0.23 to 0.50. Results did not demonstrate heterogeneity or publication bias (I2=0%). Fourteen studies investigated the role of combined modalities on the incidence of DVT. These showed a reduction in DVT from 6.18% (200/3238) in the control group to 2.05% (63/3074) in the treatment group. Odds ratio was 0.31, 95% CI 0.23 to 0.43. Results were consistent with significant heterogeneity (I2=53.5%).

Conclusions: Combined prophylactic modalities decrease significantly the incidence of venous thromboembolism, both DVT and PE compared to single modalities. The results of the current review support their use, especially in high risk patients.



P-8 Evaluation of Venous Thromboembolism Prophylaxis In Randomly Selected Medical and Surgical Patients

B. K. Zierler, J. Lee, G. Han, H. Oh, C. Jacobson - University of Washington, Seattle, WA

Background: The evidence to support the use of pharmacologic prophylaxis to prevent venous thromboembolism (VTE) has been available for years, yet VTE remains a problem in hospitalized patients. The purpose of this study was to evaluate the utilization of VTE prophylaxis in randomly selected 200 medical and surgical adult inpatients prior to the implementation of a VTE Safety Toolkit.

Methods: This study was a part of a larger pre/post study to implement a VTE Safety Toolkit that consists of clinical algorithms for the prevention, diagnosis, and management of VTE. We randomly selected 100 medical inpatients and 100 surgical inpatients at an academic medical center. Retrospective medical chart reviews of those 200 inpatients were conducted during the pre-intervention period.

Results: Of the 100 randomized medical inpatients, 47 were placed on anticoagulation prophylaxis only or anticoagulation and mechanical prophylaxis. Of these 47, 13 were on chronic warfarin therapy which means that they were already anticoagulated and therefore, did not need further prophylaxis to prevent VTE. Seven patients were ineligible for anticoagulation due to a contraindications, thus, 47 of 93 (51%) eligible patients received appropriate therapy, whereas 46 of 93 patients (49%) were not appropriately anticoagulated to prevent VTE. Only 49 surgical inpatients received pharmacological prophylaxis either anticoagulation alone or anticoagulation with mechanical prophylaxis. Eight of 28 patients (29%) were considered to be at moderate risk for VTE, 18 of 38 patients (47%) were categorized to be at high risk and 23 of 30 patients (77%) at the highest risk received pharmacological prophylaxis. Five were ineligible for prophylactic anticoagulants due to documented contraindications, such as bleeding and coagulopathies. Of the 51 surgical inpatients who did not receive pharmacological prophylaxis, 42 patients (82%) received mechanical prophylaxis. Nine patients who had major surgery did not receive any form of VTE prophylaxis.

Conclusions: Using the American College of Chest Physicians recommendations for VTE prophylaxis as the gold standard we reviewed 200 cases to determine VTE risk categories, eligibility for prophylaxis, and contraindications to anticoagulation. Approximately half of all eligible medical and surgical inpatients did not receive the appropriate protection against VTE. The dosing and timing of anticoagulation prophylaxis needs to be determined in future studies to explain the high number of VTE cases found in the patients that had been placed on anticoagulants. The VTE Safety Toolkit will be implemented in October 2007 and data will be collected 6 months following the implementation. Our goal is to increase the number of patients who are assessed for VTE and appropriately prophylaxed using institutional guidelines and organizational supports.

P-9 An Algorithm For Outpatient Deep Venous Thrombosis Management and Severe Post-Thrombotic Syndrome At Mid-Term Follow-Up

E. Sivrikoz, M. Kurtoglu - Dept. of General Surgery, Istanbul School of Medicine, Istanbul University, Istanbul, Turkey

Background: An outpatient treatment algorithm according to all etiology groups of deep venous thrombosis (DVT) was adapted. Evaluation of initial DVT treatment as well as mid-term follow-up results of severe post-thrombotic syndrome (PTS) after mean two years according to CEAP classification is reported.

Methods: Patients with acute, symptomatic, lower extremity DVT were sub-grouped into transient risk and hypercoagulability due to the designed algorithm. Initial treatment: after diagnosis a subcutaneous low-molecular-weight heparin (LMWH) (enoxaparine sodium 150 IU/kg/day), on second day an oral anticoagulant (Warfarine Na 5mg/day). Patients were equipped with Class-II below-knee compression stockings (Sigvaris-212) and sent home. Follow-up visits: 1, 3, 6 and 12th month. CEAP classification was assessed to determine chronic venous disease and severe PTS at late follow-up.

Results: Records of 121 acute DVT cases treated according to the algorithm at 2000 - 2004 were analysed. There were 3 recurrences, 1 major bleeding, 1 minor bleeding in transient risk group (n=71); 8 recurrences, 4 major bleeding, 3 minor bleeding and 3 pulmonary emboli in hypercoagulability group (n=50) after 12 months of initial treatment (Table 1). There was a significant difference regarding recurrent DVT (0.02), hypercoagulability group was associated with a three-fold increased risk for recurrent DVT (RR=3.79, 95% CI 12.30-159.17). CEAP classification was assessed on 10 patients from transient risk and 30 patients from hypercoagulability group (mean 25 ± 12.22 months) in mid-term follow-up (Table 2). Severe PTS (C- 4s, 5s, 6s) developed in 4 patients (10 %); all belonged to hypercoagulability group and all had a recurrent DVT with inadequate INR on presentation during study interval.

Conclusions: Thrombophilia investigation using an algorithm is a helpful adjunct to sub-classify patients, as well as guide therapy. Hypercoagulability is associated with a three-fold increased risk for recurrent DVT. CEAP outcomes after two years indicate relatively good PTS outcomes. Severe PTS was associated with hypercoagulability and recurrent DVT.

TABLES

Table 1. Results of initial treatment (n=121)					
	Transient risk group	Hypercoagulability Risk Group			
		Weak thrombophilia group	Idiopathic group	Strong thrombophilia group	
	(n=71)	(n=20)	(n=10)	(n=20)	
Recurrent DVT	3	2	1	5	
P.E.	0	1	0	2	
Major bleeding	1	1	1	2	
Minor bleeding	0	1	1	1	

Table 2. CEAP classification - subgroups of all etiologies					
	Transient Risk (n=10)	Hypercoagulability Risk (n=30)			
		Weak Thrombophilia (n= 10)	Idiopathic (n=5)	Strong Thrombophilia (n=15)	
C-0as	3	2	1	2	
C-0s	1		2	2	
C-1as		1			
C-1s		1			
C-2as	1	2			
C-2s	3			1	
C-3as		2		2	
C-3s	2	2		5	
C-4as				1	
C-4s			1	1	
C-5s			1		
C-6s				1	

P-10 Prevalence of Isolated (C2) and Complicated (C2+) Varicose Veins Among Patients Consulting Vascular Specialists For Varicosis: A Snapshot

M. Cazaubon¹, M. Lugli², P. Burseta³, M. Perrin⁴, F. A. Allaert, V⁵ - ¹American hospital, Neuilly, France; ²Vascular hospital, Bologna, Italy; ³vascular dpt hopital, Modena, Italy; ⁴Vascular dpt hospital, Lyon, France, ⁵cenbiotech ceren ESC and DIM CHRU Dijon, Dijon, France

Background: The revised CEAP classification allowed to differentiate isolated varicose veins (C2) and complicated varicose veins (C2,3, C2,3,4 or C2,4 or C2,3,4,5¹/₄). The main objective of this study was to identify with this revised classification the prevalence of isolated varicose veins (C2) and complicated (C2+) among patients consulting vascular specialists and to compare their symptomatic expression and the data issued from the detailed CEAP concerning Anatomy (A), Etiology (E), Physiopathology (P).

Methods: European cross sectional study conducted in euroepan countries and involving ten study centres per countries. Each centre (Phlebologist, angiologist or vascular surgeons) were asked to include the next 10 patients consulting for varicose veins and to achieve a full completion of the clinical and duplex examination required by the advanced CEAP for each leg. Exclusion critera were patients with CVD without any varicose veins and patients with acute deep venous thrombosis or superficial thrombophlebitis. Venous reflux was define by a retrograde flow in the reverse direction to physiological flow lasting for more than 0,5 seconds in the superficial veins, except the femoropopliteal veins, where the cut off was set at 1.0 second.

Results: Current preliminary results covers 171 patients (100 French and 71 Italians) presenting 258 legs with varicose veins. They were 57 \pm 15 years old, 71.6% were female and 36,5% are overweighted. C2 prevalence was 64,0% and C2+: 36,0% and strictly comparable between Italy (C2=63,3%) and France (C2=64,4%). Prevalence of C2 significantly differs between male and female : 49,3% vs 69,0% (p<0.001) and decreases according age (p<0.001). C2,C3 represents 11,2% of patients, C2,4 : 9,7%, C2,3,4 : 8,9%, C2,3,4,5: 1,9% , C2,4,5 : 1,6% and those involving C6 : 2,4%. C2+ are more symptomatic than C2 : 74,2% vs 52,1% (p<0.0005). These previous results are similar in Italy and France. A secondary etiology is significantly more frequent in C2+ (17,2% vs 1,2% p<0.0001), deep vein are more often involved (16,1% vs 1,2% p<0.001) and the association of reflux and obstruction is more present (13,6% vs 1,9% p<0.0). After duplex examination, whatever the country, the average duration to fill in the grid of the advanced CEAP is 6,1 ± 3,8 mn.

Conclusions: Our study support the usefulness of the advanced CEAP by providing a description which better reflect the real venous status of patients which allows better comparison between epidemiological studies and its usability as ascending severity classes than in the previous CEAP version.

P-11 Prospective Randomized Efficacy of Ultrasound-Guided Foam Sclerotherapy Compared To Ultrasound-Guided Liquid Sclerotherapy In the Treatment of Symptomatic Venous Malformations

T. Yamaki¹, M. Nozaki¹, H. Sakurai¹, M. Takeuchi², K. Soejima³, T. Kono¹ - 1Tokyo Women's Medical University, Tokyo, Japan; ²Nihon University, Tokyo, Japan; ³Tokyo Metropolitan Hiroo General Hospital, Tokyo, Japan

Background: To compare the clinical outcome between ultrasound-guided foam sclerotherapy (UGFS) and ultrasound-guided liquid form sclerotherapy (UGLS) in patients with congenital venous malformations (CVM).

Methods: Eighty-nine patients with symptomatic CVM were treated with ultrasound-guided scleroherapy. There were 22 males and 67 females with mean age of 14.5 years. The sclerosing agents used were 1% polidocanol (POL) or 10% ethanolamine oleate (EO). POL was injected predominantly into smaller, superficial lesions, whereas EO was used for large, deeper lesions. Foam sclerosing solution was provided using Tessari's method.

Post-sclerotherapy surveillance was done at 6 months after last session using duplex ultrasound. Findings obtained by duplex scanning were divided into 4 groups:

Disappeared: The venous space was occluded and was totally shrunk

Partially recanalized: The venous space was partially recanalized and was partially shrunk

Totally recanalized: The venous space was totally recanalized and returned at the same sizeWorsened: The venous space was totally recanalized and became worsened.

Results: Forty-nine patients were treated with UGFS and remaining 40 were treated with UGLS. The most common location of VM in UGFS was head and neck region, followed by upper extremity and lower extremity region (51%, 19%, and 14%, respectively). Similarly, in the UGLS group, head and neck was involved in 42% of the patients, 35% had lower extremity and 10% had upper extremity involvement. The amount of POL was significantly smaller in patients who were treated with UGFS (p=0.022). Similarly, there was a significant reduction in the use of EO in patients treated with UGFS (p=0.005). The proportion of CVM with total disappearance and partial recanalization was significantly higher in patients treated with UGFS (p=0.002). No major complications related to sclerotherapy were encountered in both groups.

Conclusions: These findings suggest that UGFS could have greater promise compared with UGLS in the treatment of CVM.

P-12 Elastic Stockings and Ulcer Treatment: What About Pressure and Stiffness?

G. Mosti - Clinica MD Barbantini, Lucca (LU), Italy

Background: Multilayer bandages are the basic method of choice for treating venous leg ulcers. They exert a high standing pressure (necessary to counteract venous hypertension) starting from a low and comfortable supine pressure: this high difference between standing and supine pressure corresponds to a high stiffness. Elastic stockings produce lower pressure and stiffness when compared with multilayer bandages and have a lower recommendation grade for ulcer treatment.

Aim: To compare maintenance of pressure over time and elastic properties of three compression devices by in vivo-measurements in order to evaluate an optimal compression system for the treatment of venous leg ulcers.

Methods: In 12 healthy volunteers two kinds of stockings were tested: a two-layer Gloriamed ulcer Kit®, consisting of a liner (24 mm Hg) and a Gloria 261® natural rubber stocking (23-32 mmHg), and a single Gloria 261®. Interface pressure was measured in supine and standing position and the Static Stiffness Index (SSI) was calculated by subtracting the supine from the standing pressure. Thereafter the volunteers were asked to wear the ulcer kit for one week taking off the outer layer over night. The pressure and stiffness data were compared with those measured in 12 patients suffering from venous leg ulcers and treated by means of an inelastic bandage system (Rosidal Sys®) with high pressure and stiffness.

Results: The mean pressure values (mm Hg) in the supine and standing position were 28.3 ± 4.0 and 35.3 ± 5.4 for the stocking, 45.8 ± 5.8 and 55.2 ± 5.1 for the ulcer kit, 69.5 ± 5.8 and 94.8 ± 10 for the bandage sytem. Differences are statistically significant between all the compression devices.

The corresponding SSI values were 6.9 ± 2.4 (stocking), 9.3 ± 4 (ulcer kit) and 25.3 ± 6.1 (Rosidal Sys ®). After 48-72 hours the pressure loss in the supine and standing position was 6.1% and 5.4% with the ulcer kit, but 42.1% and 36,1% with the bandage system (p<0.001) so that the pressure range of the bandage system came close to that of the ulcer-kit.

Conclusions: The ulcer-kit achieves a standing and working pressure similar to that exerted by multilayer bandages, at least after some days when there is a considerable pressure loss of the bandage system. Since the outer layer is removed over night the resting pressure of the liner keeping the ulcer dressing in place is well tolerated.

This new stocking system can be recommended at least for "not complicated" ulcers with a surface less than 100 cm2 and lasting less than 1 year.

P-13 Inelastic Compression Increases Venous Ejection Fraction More Than Elastic Bandages

G. Mosti¹, H. Partsch², V. Mattaliano¹ - ¹Clinica MD Barbantini, Lucca (LU), Italy; ²Dermatology Department, Vienna, Austria

Background: One of the pathophysiological key-parameters of an incompetent venous pump is the reduced ejection fraction (EF) from the lower leg. The aim of our work was to investigate if compression therapy is able to increase EF in patients with chronic venous insufficiency and if there is any difference between elastic and inelastic bandages.

Methods: EF was measured by means of strain gauge plethysmography in 14 healthy subjects (coefficient of variation $6.5\% \pm 2.4$) and in 20 patients (CEAP C2-C6) waiting for venous surgery. The probe was placed 5 cm distal to the patella in supine position. After calibration, the examined leg was elevated to empty the calf veins and then the patient was asked to stand up and wait until a stable signal was achieved. The resulting volume increase after refilling of the veins is defined as venous volume (VV). Then the patient was asked to perform 20 standardized steps in 20 seconds. The resulting volume decrease corresponds to the ejected volume (EV) and ejection fraction is calculated according to the formula EF= EV/VV x 100. The procedure was repeated with an elastic and an inelastic bandage on the leg applied with the same resting pressure distally to the strain gauge. The interface pressure of each bandage was measured about 12 cm above the inner ankle using an air-filled pressure transducer in supine and standing position.

Results: With the same resting pressure of about 40 mm Hg , the standing pressure raised by 4.9 ± 2.0 mm Hg with an elastic and by 18.7 ± 4.3 mm Hg (P<.001) with an inelastic bandage. EF increased from $36.4\% \pm 10.2$ (baseline) to $48.2\% \pm 10.7$ with the elastic and to $62.3\% \pm 8.1$ (P<.001) with the inelastic bandage, coming close to the normal range ($63.8\% \pm 2.4$). Standing pressure and the difference between systolic and diastolic pressure during exercise, both being higher with inelastic bandages, showed a significant correlation with EF (r= .58 and r= .65 respectively).

Conclusions: Ejection fraction which is severely reduced in venous insufficiency can be increased by compression therapy. Inelastic compression is much more effective than elastic bandages and is able nearly to normalize ejection fraction.

P-14 Arterial Revascularization and Compression Therapy In the Treatment of Mixed Arterial/Venous Leg Ulcers

D. J. Milic, S. S. Zivic, Z. D. Perisic, R. Jankovic, G. Djordjevic, D.M. Stamenkovic, Z.D. Maksimovic Clinic for Vascular Surgery, Clinical Centre Nis, Serbia

Background: Venous ulcers are a major health problem because of their high prevalence and associated high cost of care. The natural history of this disorder is slow healing and high recurrence rate. About 3%-5% of patients with venous ulcers have combined arterial and venous insufficiency. The aim of this study was to show the results in the treatment of mixed arterial/ venous ulcers with and without compression therapy after performing arterial revascularization procedures.

Methods: A total of 27 patients (11 women and 16 men; mean age 65.7 years) with mixed arterial/venous ulcers were randomized into two groups: treatment group (14 patients) and control group (13 patients). All 27 patients had an active ulcer in the gaiter area (ulceration surface: 10-44 cm2; duration: 5 months - 4 years), deep vein reflux confirmed with color duplex scan and ankle brachial pressure index of 0.7 or less. Mean ankle brachial pressure index was preoperatively 0.53 and 0.49 in patients in the treatment group and 0.50 and 0.52 in patients in the control group). Arterial revascularization procedures were performed in all 27 patients included in the study (7 aortobifemoral, 12 above-knee femoropopliteal, 4 belowknee femoropopliteal and 4 femorotibial reconstructions). Thirty days after operation patients in the treatment group underwent compression treatment using multi-layer bandaging system while patients in the control group didn't receive any additional treatment. The results in the treatment of ulcers in these two groups were analyzed in terms of healing rate, time for healing and recurrence rate during the one year follow-up period. Primary endpoint of the study was complete ulcer healing at 24 weeks. After ulcer healing, patients in the treatment group wear compression stockings class II (25mm Hg) in order to avoid recurrence. The study excluded patients with heart insufficiency (EF<35), pregnancy, cancer, rheumatoid disease and diabetes. Patients with foot or/and finger ulcers were also excluded from the study.

Results: Postoperative mean ankle brachial pressure index was 0.85 and 0.84 in patients in the treatment group and 0.86 and 0.92 in patients in the control group. The healing rate was 71.4% (10/14) in the treatment group, and 38.5% (5/13) in the control group (p<0.01). Median ulcer healing time was 68 days (27-141 days) in the treatment group versus 89 days (41-146 days) in the control group (p<0.05). The recurrence rate during the one year follow up period was 40% (4/10) in the treatment group and 40% (2/5) in the control group.

Conclusions: The study suggests that mixed arterial/venous ulcers could be successfully treated with compression therapy after performing arterial reconstructive procedures.

P-15 Morphological Changes On Varicose Vein Wall Corresponds To MMP/TIMP Alterations

B. Aravind¹, T. Navin², C. Monaco², E. Paleolog², A. H. Davies¹ - ¹Imperial College, Charing Cross Hospital, London, United Kingdom, ²Kennedy Institute of Rheumatology, Imperial College, London, United Kingdom

Background: Primary weakness of vein wall is widely accepted as aetiology of varicose vein and is ascribed to its changes in matrix composition . Proteases, especially Matrix Metalloproteases (MMPs), and its inhibitors, TIMPs, are important regulators of matrix turnover. It was hypothesised that alteration in these proteases causes matrix imbalance which will correspond to observed changes in morphology of vein wall. Vein wall thickness was considered as an index of morphological change.

Methods: Stripped varicose vein were collected from patients undergoing operation as proximal (groin end) and distal (knee end) segments. The wall thickness were measured from EVG stained sections, as distance between external elastic lamina and luminal endothelium. The segments were further subdivided as hypertrophic (>1000µm, wall thickness) and atrophic (<500µm) to study the extremes of changes. Sections were stained by immunohistochemistry using antibodies to MMP-2, MT1-MMP, TIMP-2 and TIMP-3. They were image analysed using AnalySIS software to quantify the expression of proteases. Vein segments were also assayed for MT1-MMPenzyme activity.

Results: The distal varicose vein wall were thinner (n= 24 pairs, median thickness 574μ m) than proximal (766 μ m, p>0.07). Atrophic vein segments were predominantly from the distal end of varicose veins (38% to 8% in proximal).

TIMP-2 and TIMP-3 expression were increased in the thicker proximal segments compared to distal (4.34 versus 1.29% per hpf, p>0.06 and 0.94 versus 0.41%, p>0.39, respectively). In comparison, MT1-MMP activity was significantly increased in distal segments (2.78 versus 0.72pg/ μ m in proximal, p<0.04). TIMP-2 and TIMP-3 expressions were higher in hypertrophic segments compared to atrophic segments (4.34 versus 0.99%, p>0.05 and 1.65% versus 0.08%, p>0.10 respectively). The reverse was true of MT1-MMP (1.05 versus 1.90, p>0.97) and MMP-2 (0.95 versus 1.09%, p>0.85), both showing a higher expression in atrophic segments.

Conclusions: This internal comparison of proximal and distal varicose vein wall highlighted the link between vein wall thickness and expression of MMP and TIMP. The proximal vein wall which was found to be thicker, has a higher expression of protease inhibitors, TIMP-2 and TIMP-3, which inhibits matrix turnover. This favours deposition of matrix components like collagen, resulting in thickening of the vein wall. The reverse was true in distal segments, where a higher activity of protease MT1-MMP activity increases the matrix turnover, reducing matrix content and leading to thinning of the vein wall. This relationship was again reflected in hypertrophic and atrophic segments. This study was able to prove conclusively the link between altered protease expression and morphological changes. The focus of future studies should now shift to the possible triggers for such changes in varicose vein wall.

P-16 Lower Power Improves Clinical Outcome of the Endovenous Laser Treatment

S. Kaspar, J. Siller, Z. Cervinkova - Flebocentrum, Hradec Kralove, Czech Republic

Background: Many controversies still remain as to best parameters of the endovenous laser treatment of varicose veins and to date there is no standardized energy delivery protocol. Based on our experimental laboratory study we performed retroprospective clinical study of our patients operated on with endovenous laser during 4.5 year period.

Materials And Methods: 463 procedures in 430 patients were analysed. Post-operative follow-up was accomplished after 1 month, 6 months and yearly thereafter.

Cox regression analysis was used to detect factors influencing non-occlusion and early or late recanalisation of saphenous vein. Results were evaluated by comparison of CEAP clinical class pre- and post-operatively, by percentage of recanalizations and using Kaplan-Meier life-table method. Postoperative data were available during different time periods in 457 limbs (98.7%).

Results: Saphenous occlusion was verified in 446 limbs (96.3%) after 1 month, non-occlusion or early reopening was seen in 17 limbs (3.7%) at this time. Totally, 40 non-occluded saphenous trunk veins were found during the whole follow-up period (1-54 months) which represents final occlusion rate of 91.36%. Using Kaplan-Meier analysis, we reached 83% occlusion rate during follow-up period up to 4.5 years. Mean clinical CEAP classification improved from 2.22 (before operation) to .24 (1 month after) and .48 (last visit).

Cox regression analysis selected 2 factors with statistical significance: body mass index (p = .017) and laser power (p = .031).

Cumulative rate of occlusions in 54 months horizon is significantly higher (86%) in patients with BMI < 25 compared to patients with overweight (63%), p = .00032.

When comparing the influence of laser power on quality of saphenous occlusion, treshold of 13W was set arbitrary based on median values in occluded and non-occluded cohorts and using Kaplan-Meier survival method, the results of treatment with power < 13W and > 13W were analysed. Using power values < 13W, results were significantly better (logrank test: p = .048, Cox-Mantel test .02) compared to power values of 13 W or more. Median power in non-occluded veins was 14W while in occluded trunks 13W. This difference is statistically significant (p = .0095).

Conclusions: Present clinical study supports concept of "slow and gentle heating" during the endovenous diode laser treatment of varicose veins to achieve good immediate and late result. Based on our observations and their statistical analysis, we recommend lower or medium power settings (8 to 13 W) with slower pull-back speed of laser fibre (0.2 to 2 mm/sec) to achieve sufficient energy per centimeter of the vein and the optimal clinical outcome with minimal side-effects.

P-17 Effectiveness of Weight Loss On the Evolution of Chronic Venous Insufficiency (CVI) After Bariatric Surgery In Obese Patients

J. Benigni¹, J. Uh^p, J. Gobin³, A. Capella⁴ - ¹Hôpital BEGIN, St Mandé, France; ²Surgical Venous Center, Neuilly, France; ³Vascular medecine, Lyon, France; ⁴Angiologist, Paris, France

Background: In patients with morbid obesity, surgical treatment is more effective than non surgical treatment for weight loss and control of some comorbid conditions (Maggard 2005).

CVI is a known complication in these patients and 2/3 of them have no venous reflux (Padberg 2003). Bariatric surgery (BS) corrects the complications of CVI in almost all patients with a Body Mass Index (BMI) > 40 (Sugarman 2001). The objective is to study the relationship between weight loss and CVI in obese patients (BMI > 30) after BS.

Methods: From a database of 758 obese patients examined before a BS (clinical and duplex ultrasound examination), patients with CVI (Ceap C3-C6) and a BMI > 30 were reviewed.

Results: Of the 758 patients, 57 (7.5%) met the criteria, 3 did not have a BS, 1 refused to be reviewed. 35 who had a BS, were reviewed (65%). Of the 35 reviewed patients : 30 were females and 5 males. Mean age was 45.74 +/-11 years. The distribution of CEAP clinical class was C6 (n=0), C5 (n= 1), C4a and b (n= 4 and 2), C3 (n= 28). A venous reflux was present in 36% patients. 60% wore a compression for one month. Mean BMI was before BS 47.72 +/- 9.05 and after BS 41.33 +/- 10. Mean weight loss was 32.13 +/- 17 kg. Mean period between BS and the review was 16.35 +/- 9 months. After BS, the outcome parameters were based on clinical signs : group I- no clinical improvement/aggravation, group II - clinical improvement (decrease of oedema, lipodermatosclerosis, or pigmentation), group III- disappearance of clinical signs. The distribution was : group I n = 5 patients, group II n = 13 patients, group III n = 17 patients Mean weight losses (kg) were in group I : 3 +/-7.1, group II : 34.9 +/- 3.9, group III : 36.9 +/- 3.5

There was a significant correlation between the importance of weight loss and the improvement of CVI (t-test of Student). The difference was very significant between the groups II and III and the group I (p < 0.0006). No difference on CVI evolution was found between the groups with and without compression.

A possible bias : 19 lost to follow up patients but there was no difference between the lost to follow up group and the reviewed group for BMI, weight, sex, age, distribution of CEAP and venous reflux.

Conclusion: In obese patients with CVI, weight loss after BS could be a major parameter of improvement of clinical signs of CVI.

P-18 One Year Follow-Up of Radiofrequency Segmental Thermal Ablation (RTFA) of Great Saphenous Veins

T. M. Proebstle¹, B. Vago², J. ALm³, O. Goeckeritz4, C. Lebard⁵, O. Pichotó - 1University of Mainz, Mainz, Germany; ²University of Heidelberg, Heidelberg, Germany; ³Dermatoligicum, Hamburg, Germany; ⁴Venenzentrum am Elsterpark, Leipzig, Germany; ⁵Hospital St. Michel, Paris, France; ⁶CHU Service de Chirurgie Vasculaire, Grenoble, France

Background: Radiofrequency segmental thermal ablation has been introduced recently and shown its feasibility in occlusion of incompetent great saphenous veins (GSVs).

Methods: N = 295 GSVs in 225 patients were treated by RTFA under local anesthesia in a prospective multicenter trial. Duplex-ultrasound control visits were performed after 3 days, 3, 6 and 12 months. Clinical data was obtained at the same time

Results: Of 225 patients n=166 (73.8%) were female with a mean age of 50.5 years [range 18 - 79]. All GSVs were treated per protocol with double cycle treatment at the first segment with an average length treated of 36.9 cm. Concomitantly performed procedures were phlebectomy in 164 legs (55.6%) and sclerotherapy in 38 legs (12.8%). During followup, one vessel was open at 3 days but occluded thereafter. Treatment failures or recanalizations of the once occluded GSVs were n = 1 of 284 at 3 months, n = 2 of 250 at 6 months and n = 1 of 111 at 12 months after the intervention. The corresponding occlusion rates calculated according to the method of Kaplan and Meier were 100%, 99.6%, 98.9% and 98.0%, respectively. Of n = 111 GSVs, which were followed for 1 year, inner diameters were measured with ultrasound 3cm distal to the saphenofemoral junction (SFJ). N = 54 of these GSVs were sonographically not detectable after 1 year. Of the remaining n = 57 GSVs with a complete data set, the average diameter at 3 cm distal to the SFJ reduced from 5.3 \pm 1.8 mm pre treatment to 4.3 \pm 1.7 mm at 3 days, to 2.9 \pm 1.0 mm at 6 months and to 2.4 ± 1.1 mm at 1 year after treatment. The average VCSS score improved from 3.9 ± 2.1 before treatment to 3.6 ± 1.2 at 3 days, to 1.1 ± 1.7 at 3 months, 0.4 ± 0.8 at 6 months and 0.5 ± 1.2 at 12 months thereafter. Presence of any pain in the treated limb improved from 58.6% before treatment to 25.2 %, 7.3%, 5.2% and 7.2% at the same follow-up intervals. Likewise, presence of leg swelling improved from 52.9% to 5.4%, 8.8%, 7.2% and 2.7%. Side effects noticed at any time during follow-up in the RSTA-treated area were ecchymosis (5.8%), paresthesia (3.4%), erythema (2.0%, skin pigmentation (2.0%), hematoma (1.4%) and phlebitis (1.0%).

Conclusion: RSTA showed a high success rate and durability of the once achieved occlusion of treated GSVs together with a moderate side-effect profile. A remarkable subsequent improvement of clinical symptoms was noted.

P-19 Elimination of Superficial Reflux With Or Without Subcutaneous Fasciotomy - The Impact On Deep Axial Reflux and ulcer Healing

J. T. Christenson - Division of Caraliovascular Surgery,University Hospital of Geneva, Geneva, Switzerland

Background: Deep venous reflux (DAVR), without history or Duplex ultrasound evidence of previous deep vein thrombosis (DVT) can be seen in patients with greater saphenous reflux. DAVR is suggested an important contributor to skin changes and venous ulcer. Saphenous vein ablation eliminates deep vein reflux in 1/3 of patients with DAVR. The explanation of this phenomenon is still unclear. Tissue pressures (intramuscular, Pim and subcutaneous, Psc) are increased in C6 patients. Saphenous vein ablation alone decreases Psc tissue pressure, while additional subcutaneous fasciotomy also lowers Pim resulting in improved ulcer healing.

Patients and Methods: This study included 25 limbs in 22 patients. Inclusion criteria: Severe venous insufficiency therapy resistant venous ulcers planned for surgical eradication of superficial reflux with or without subcutaneous fasciotomy. Exclusion criteria: History of DVT, previous venous or limb surgery, lesser saphenous vein reflux or deep vein thickening or distortion on Duplex ultrasound scanning.

Patients were assigned to either of 4 subgroups for purpose of analysis; DAVR with fasciotomy (Group1, 5 limbs), DAVR without additional fasciotomy (Group2 5 limbs), no DAVR with fasciotomy (Group 3, 10 limbs) and no DAVR without additional fasciotomy (Group 4, 5 limbs). All patient characteristics were comparable between the groups. Tissue pressure measurements were performed pre- and postoperatively and at 3 months. Ulcer healing was monitored. Changes in deep segmental and axial reflux were compared preoperatively and at 3 months, and the impact of additional subcutaneous fasciotomy and deep axial reflux evaluated.

Results: DAVR was eliminated in 2/10 limbs (20%) following surgery. When additional fasciotomy was performed axial reflux was eliminated in 2/5 limbs (40%), compared to controls (no fasciotomy), 0/5 (0%). Elimination of segmental reflux was higher in patients who had additional fasciotomy (p=0.0174) compared to controls.

Intramuscular tissue pressures in limbs evaluated (mmHg) and ulcer healing:

		Preop.	p- value	Postop.	p- value	3 months	Ulcer healing at 3 months
	Fasciotomy	28.4±2.7	<0.001	8.8±2.3	0.005	4.8±1.9	80%, 4/5
Axial Reflux	p-value	n.s		<0.001		<0.001	n.s.
	No fasciotomy	25.8±4.0	<0.001	18.8±4.7	<0.001	11.0±4.2	40%, 2/5
	Fasciotomy	18.1±3.1	<0.001	4.4±2.0	n.s	4.5±1.8	90% 9/10
No axial reflux	p-value	n.s		<0.001		<0.001	0.049
	No fasciotomy	18.2±2.2	n.s.	19.8±3.9	n.s.	19.4±2.5	20%, 1/5

Presence of DAVR increases tissue pressures. In patients with DVAR having additional fasciotomy had higher postoperative Pim than patients with absence of DAVR, while patients with DAVR and no fasciotomy had decreasing Pim at 3 months in limbs where DAVR was resolve.

Conclusion: Venous ulcer disease is linked to venous hypertension, but also increased tissue pressures. Once pressures are decreased, following elimination of superficial reflux and subcutaneous fasciotomy, improved ulcer healing occurs. When primary DAVR is present intramuscular tissue pressure is higher compared to case with no DAVR. Additional fasciotomy lowers Pim over time as DAVR is eliminated.

P-20 The Effectiveness and Use of Compression Stockings of Various Strength For the Treatment of Venous Disorders and Diseases: A Literature Survey

W. Blaettler¹, H. E. Gerlach², F. Amsler³ - ¹Angio Bellaria, Zürich, Switzerland; ²Center for Vascular Diseases, Mannheim, Germany; ³Amsler Consulting, Biel-Benken, Switzerland

BACKGROUND: The strength of medical compression stockings (MCS) optimal for the treatment of the various manifestations of venous disorders (CVD) and insufficiency (CVI) is not known.

METHODS: We reviewed the literature for RCTs and large registries on leg compression for occupational symptoms and oedema (CVD, C0s-C3s), symptoms of acute DVT, prevention of the PTS, and chronic venous leg ulcers (C6). Studies were included if they compared one MCS with another one of different strength and/or with no compression or bandaging.

RESULTS: Eleven trials on CVD covered 1.453 subjects. MCS exerting an ankle pressure of 10-20mmHg had a clear effect on symptoms and oedema as compared with <10mmHg, placebo stockings, or no treatment (p20mmHg.

The symptoms of acute DVT were investigated in 2 comparable studies (n=81). MCS 20-30mmHg, 30-40mmHg and bandages provided better relief than bed rest (p<.01) with no difference between the compression modalities.

The prophylaxis of DVT was explored in 2 studies of identical design (n=374). MCS 30-40mmHg were started as symptoms vanished and prevented about half of the PTS as compared with no treatment (p<.01).

Three consecutive German registries (n=2.149) investigated the real world practice. Leg compression was started immediately upon diagnosis of DVT in 98, 99 and 93%, respectively. In the latest trial, bandages were applied initially in 38% and MCS in 62%. MCS 20-30mmHg were used in 93%.

Three studies on ulcer healing at 90 days (n=134) revealed a better healing rate of MCS 20-30mmHg than bandages (65 versus 80%, p<.02). Two trials using a similar plan (299 patients) found specifically designed 30-40mmHg MCS not better than bandaging (47 versus 54%).

CONCLUSIONS: In the clinical situations covered by this survey direct or indirect comparisons revealed that MCS of lower strength were at least as effective as the stronger ones or bandages. The recommendation ensuing from the RCTs on PTS prevention, i.e. to prescribe 30-40mmHg MCS, is not followed by the vast majority of German vascular physicians who use 20-30mmHg MCS. Confirmation of the observed trend in further direct comparison studies would render compression therapy more acceptable without loosing effectiveness.

P-21 Management of Venous Injuries At the Air Force Theater Hospital In Balad, Iraq

> S. Gifford¹, W. T. Jones¹, M. A. Ricci², W. D. Clouse¹, T. E. Rasmussen¹ - ¹Wilford Hall Medical Center, San Antonio, TX; ²University of Vermont, Burlington, VT

Background: Wartime venous injuries may present life-threatening and challenging treatment problems. In order to better understand the nature of venous injuries, we have reviewed the experience of the Air Force Theater Hospital (AFTH) during Operation Iraqi Freedom.

Methods: All vascular injuries treated at the AFTH were registered from September 1, 2004 until August 31, 2007. Data recorded included basic information about the injury and treatment.

Results: Over the period of study 583 cases were entered into the registry. Of these, 226 (38.8%) had venous injuries, of which 47 had an isolated venous injury (20.8%). The mechanism of injury was an explosive device in 115 cases, a gunshot wound in 98, blunt trauma in 4, iatrogenic trauma in 2, and a dog bite in 1 case with the remainder (6) by unknown mechanisms. The site of injury was the lower extremity in 130 cases, the upper extremity in 42, abdomen in 31, neck in 27, and chest in 12. Twenty-three had more than one anatomic area injured. In the majority of cases, ligation was the preferred treatment (103 or 45.6%). Lateral venoraphy was used in 72 cases (31.9%) and venous bypass was used in 26 cases (11.5%). End-to-end anastomosis, panel graft, and shunt were also utilized in only one case each. Eight patients (3.6%) died of their combined injuries with one week of hospitalization.

Conclusions: Wartimes venous injuries are less frequent than arterial injuries. Both the variety of explosive devices encountered as well gunshot wounds are likely to produce venous injury which is most likely to occur in the extremities. Repair (venoraphy or bypass) were utilized with roughly the same frequency as ligation but further follow up on the long term outcome is needed.

P-22 Critical Issues In the Management of Venous Malformation (VM) Coexisting With Lymphatic Malformation (LM) - Klippel Trenaunay Syndrome (KTS)

B. Lee, J. Laredo, D. Deaton, R. Neville - Georgetown University, Washington, DC

Background: Klippel Trenaunay Syndrome (KTS) has been known for the complexity of interwound hemodynamics among its various vascular malformation components. The 'marginal vein' as venous malformation (VM) component of KTS has been mandated to reduce the risk of chronic venous insufficiency (CVI) and pulmonary embolism (PE). But its treatment impact to coexisting lymphatic malformation (LM) component has not been clearly understood.

Methods: We made a retrospective analysis on the KTS cases that underwent the marginal vein resection to assess its impact to coexisting LM.

All N=66 patients were investigated with non- to less-invasive tests: Duplex ultrasonography, MRI, and lymphoscintigraphy. Occasionally percutaneous direct puncture lymphangiography was added to confirm the extratruncular LM, which is infiltrating lymphangioma.

Among N=32 patients with marginal veins, N=27 underwent surgical excision: one stage resection on N=22 with normal deep system, and multistage resection on N=5 with borderline hypoplastic condition.

N=5 among N=32 treated conservatively due to deep vein hypoplasia/ aplasia.

Hemodynamic change by the marginal vein compression test on Duplex ultrasonography was interpreted as an increased risk of postoperative venous hypertension by overloading to hypoplastic deep system following the forced diversion of the venous flow from the marginal vein following the resection.

Among N=27 for the resection, N=17 was confirmed for coexisting truncular LM lesion, that is, primary lymphedema as only LM component of KTS; N=7 for extratruncular LM, that is, inflitrating lymphangioma. N=3 for both truncular and extratruncular LMs together.

Results: All N=27 underwent the marginal vein resection under general anesthesia successfully.

N=23 showed excellent to good hemodynamic response and did not develop postoperative acute venous hypertension either through one stage or multistage resection. But N=4 (N=2 through one stage and N=2 through multistage) developed acute venous hypertension (eg. swelling) but all subsided within one month except N=1 which took 6 months for the relief with the conservative measurement (e.g. compression bandage and anticoagulation).

Among N=10 with extratruncular LM, alone (N=7) or combined with truncular LM (N=3), N=3 developed a significant lymphatic leakage; N=2 resolved spontaneously within 3 months but N=1 were not completely resolved with recurrent bouts of leakage and local/systemic sepsis.

None with truncular LM alone (N=17) developed lymphatic leakage but N=2 combined with extratruncular LM have shown further deterioration of lymphatic function on the lymphoscintigraphy although clinically stable with CDT based therapy (follow up - average 3.2 years).

Conclusions: A careful assessment of potential impact of the VM treatment to coexisting LM lesion is warranted for safe management of VM component of KTS; hemodynamic interreaction among the vascular malformation components should remain as a guideline for the therapy.

P-23 Variability of Interface Pressure Exerted By Compression Bandages and Standard Size Compression Stockings

H. Partsch¹, W. Vanscheidt² - ¹Medical University Vienna, Vienna, Austria; ²University Clinic for Dermatology, Freiburg i.Br., Germany

Background: The "dosage" of prescribed interface pressure plays an important role determining the efficacy of compression therapy in chronic venous disorders.

Aim: To measure the variability of interface pressure of short- and long-stretch bandages under different methods of application.

Methods: The interface pressure of short- and long-stretch compression bandages was measured under different methods of application (ie, low vs medium pressure, one vs two layers). The same practitioner applied the bandages to 12 legs of healthy subjects, and obtained measurements in supine and standing positions immediately following bandage application. These measurements were repeated with custom-made compression stockings fitted to the same legs.

Results: Two layers of short- and long-stretch bandages applied with low or medium pressure provided significantly (p=0.001) higher interface pressure when compared with one-layer bandages. When the coefficient of variation was calculated for each set of conditions, it was found that significantly higher coefficients (indicating higher variability) were achieved with bandages applied with one layer (p=0.05), or low pressure (p=0.01). Interface pressure for a custom-made Class II compression stocking at four measuring points in 12 legs, both in supine and standing positions, also demonstrated considerable variability. The coefficient of variation of the compression stocking as measured at medial distal lower leg was lower when compared with the coefficient of variation of the bandaging systems. (16.7, 18.9 and 24.5%, in standing position for the stocking, short stretch and long stretch bandages respectively, applied with medium pressure).

Conclusion: The wide range of pressure that was observed suggests that many clinical applications of compression bandages and commercially available compression stockings may not provide the intended pressure.

P-24 Recanalization of Short Saphenous Vein After EVLT S. Shokoku - Varix Ambulatory Surgery Center, Okayama

Daiichi Hospital, Okayama-shi, Japan

Background: Recanalization after endovenous laser treatment (EVLT) is one of the most disappointed events. Incidence of recanalization after EVLT of SSV (10 %, 3/30) was higher than that of GSV (0.6 %, 1/175). The purpose of this study was to determine the indicative parameters reflecting the recanalization of SSV.

Methods: 30 cases of EVLT of SSV in 28 patients (25 females, 3 males; mean age 57.3 years) between April 2005 and August 2007 were reviewed. All veins were treated with 980 nm diode laser energy delivered into the SSV via a 400 or a 600 micro meter optical fiber. Tumescent anesthesia was delivered perivenously under US guidance. Patients were evaluated clinically and with duplex US at 1 week, 1 month, 3 months, 6 months, 1 year and yearly thereafter to assess treatment efficacy.

Results: Mean diameter of SSV was 6.9 mm and mean VFI was 3.5 ml/ sec. The CEAP clinical distribution was C0 1 limb, C2 22 limbs, C3 5 limbs and C4a 2 limbs. Mean follow-up period was 258.4 days (8 - 765 days). Length of treated vein was 19.8 cm (15 - 30 cm). Average withdrawal speed was 1.7 mm/sec (1.2 - 2.8 mm/sec). Average linear endovenous energy density (LEED) was 57.5 J/cm (28.7 - 83.4 J/cm). Successful occlusion of the SSV, defined as absence of flow on color Doppler imaging, was noted in all immediately after treatment. There were three cases of recanalization at 91 days, 182 days and 467 days. In one case, main parameter influenced vein recanalization seemed the low delivered energy (28.7 J/cm). In other two cases, there were no clear differences in the delivered energy with nonrecanalization cases. The distance from SPJ to the first branch was very short and at time of recanalization, the first branch appeared again clearly. Short ablated segment of SSV was forced open by high backpressure.

Conclusions: These findings suggests a precise duplex US evaluation of branching pattern at SPJ and additional ablation of branch are needed to accomplish durable occlusion of vein lumen.

P-25 Endovenous Laser Ablation Compared With Stripping - Multi-Center RCT In Japan

T. Ogawa¹, S. Hoshino¹, S. Makimura², H. Shigematsu², N. Azuma³, T. Sasajima³, H. Sugawara⁴, M. Ichiki⁴, S. Shokoku⁵ - ¹Fukushima Daiichi Hospital, Fukushima, Japan; ²Tokyo Medical University, Tokyo, Japan; ³Asahikawa Meidical College, Asahikawa, Japan; ⁴JR Sendai Hospital, Sendai, Japan; ⁵Okayama Daiichi Hospital, Okayama, Japan

Background: Endovenous laser ablation is becoming one of the optimal treatments of saphenous varicose veins. The advantage of this procedure seems to be less invasive with good outcome compared with traditional stripping. We conducted prospective multicenter randomized comparative study of endovenous laser ablation (EA) with venous stripping.

Methods: A total of 92 patients (C2: 36 cases, C3: 35 cases, C4a: 21 cases) with 84 primary great saphenous vein (GSV) reflux, with 8 primary small saphenous vein (SSV) reflux participated in this study underwent randomly EA or stripping with the ratio of 2 EA to 1 stripping in 5 Japanese medical institutes. Stripping in 30 cases was performed from femoral junction to knee level of GSV or from popliteal junction to the end of venous reflux point of SSV under venous or general anesthesia. Endovenous ablation in 62 cases was done same part at GSV or SSV under local and tumescent anesthesia. Laser instrument was ELVeS 980nm diode laser, Biolitic, Germany. Set power was 9-12 W, average treatment energy was 43.9 J/cm. All cases wore stocking for 1 week after procedures. The results were evaluated by duplex ultrasound and air phlethysmographic examination, CVIQII QOL score and clinical examination with 6 months follow up.

Results: The complete obstruction of saphenous vein by EA was observed in 94 %. Recanalization of saphenous vein was found in 2 cases at 72 hrs after EA, 2 cases at 24 weeka after EA., There was significant improvement of QOL score and venous reflux in both EA and stripping group. The duration of hospital stay in stripping group (Average 1.5 days) was longer than that in EA group (Average 2.7 days). The frequency of adverse events after procedures was not different between two groups.

Conclusions: Endovenous laser ablation is useful for saphenous varicose veins as well as stripping. The main advantage of endovenous laser ablation over stripping was short hospital stay.

P-26 Pulse*Spray Sclerotherapy Study: A Pilot Study

J. I. Almeida, J. K. Raines - Miami Vein Center, Miami, FL

Background: Great saphenous vein (GSV) reflux is the most frequent cause of chronic venous disease. Surgical ligation and stripping of the GSV, thermal ablation with laser or radiofrequency, and sclerotherapy (chemical ablation) are the most effective treatment options. A novel endovenous catheter, which delivers a liquid sclerosant via a pulsed*spray, has received Federal Drug Administration (FDA) clearance as a delivery system. The purpose of this pilot study was to address safety and efficacy associated with this delivery system.

Methods: Over a two month period (December 2006 and January 2007), 10 subjects underwent endovenous pulsed*spray infusion of 3% sodium tetradecyl sulfate (STS) into an incompetent GSV. The range of GSV diameters was between 4 mm and 9 mm. Male or female subjects between 21 and 75 years of age were selected using standardized inclusion / exclusion criteria. The amount of sclerosant delivered was 50% of supine vein volume. All procedures were performed under local anesthesia by a single vascular surgeon. The complete protocol included eight follow-up visits to determine incidence of adverse events, degree of recanalization, and satisfaction with the procedure. These follow-up visits occured at 48-hours, 1-week, 1-month, 3-months, 6-months, 12-months, 18-months, and 24-months following the procedure. Recanalization in the treated vein segment was determined on the basis of ultrasound examinations.

Parameter	Value	Notes
Number of Veins	10	NA
Number of Subjects	8	2 bilateral procedures
Mean Age / Range (years)	51.6 / 36 - 71	NA
% Female Gender	88	By subject
Mean Treated Segment Length/ Range (cm)	35.6 / 22 - 48	Measured during surgical procedure
Mean Vein Diameter / Range (mm)	4.7 / 4.1 - 5.9	Measured by ultrasound @ time of procedure
Mean Sclerosant Volume / Range (ml)	3 / 2 - 4	Calculated based on vein diameter and length
Average Stump Length / Range (cm)	0.71 / 0.31 - 2.29	@ saphenofemoral junction
Segments of Incomplete Ablation (SIA) - %	50	Demonstrated some time during follow-up
Primary Closure (%)	80	NA
Mean Follow-Up / Range (days)	218.1 / 186 - 254	NA

Results: Results @ 6 months are summarized in the following table:

At 48-hours one subject was noted to have thrombus extension from the GSV into the common femoral vein. This was resolved without treatment at the 1-week visit. At 6-months one subject developed a small fat necrosis at the mid-thigh which required incision and drainage.

Conclusions: Pulsed*spray infusion of liquid STS resulted in 80% primary closure of the incompetent GSV. It is a quick office-based procedure requiring minimal capital expenditure. It approaches the ablative capacity of foam sclerotherapy without worrisome risks of paradoxical embolization.

P-27 Incompetent Perforators - What We Think We Know

P. A. Hertzman - Vein Care of New Mexico, Los Alamos, NM

Background: The role of incompetent perforators (IPS) in the pathogenesis of various sequelae of Chronic Venous Insufficiency and the effectiveness of ablation of these vessels using Ultrasound Guided Endovenous Chemical Ablation (UGECA) or other methods is uncertain and controversial. Due to the lack of evidence-based information, a survey was developed with the objective of establishing a consensus among experienced phlebologists related to these issues.

Methods: The survey which consisted of 14 statements using a 0-10 scale (0= strong disagreement; 10=Strong agreement) and 6 additional questions related to treatment technique, was emailed to 30 experienced phlebologists in the US, Canada, and Europe.

Results: Of the 30 surveys sent, there were 28 responses and 26 completed surveys. The Mean Scores related to the clinical significance of IPS, the effectiveness of treating IPS, and the use of Ultrasound Guided Endovenous Chemical Ablation (UGECA) for treatment are listed below:

Mean	SD	Range	
IPS Contribute to the development of:			
Lipodermatosclerosis	7.88	2.34	2-10
Venous Ulcers	8.77	1.82	2-10
Varicose Veins	7.54	2.90	2-10
IP Ablation:			
Improves Deep Venous function	6.85	2.85	0-10
Improves Lipodermatosclerosis	7.88	2.34	2-10
Improves Ulcer healing	8.65	1.60	4-10
Improves Edema	7.32	2.32	1-10
Reduces Venous Ulcer Recurrence	8.42	1.79	4-10
UGECA for IP Ablation			
Is effective	8.69	1.85	3-10
Is the optimal treatment	7.00	2.31	2-10
Is contraindicated by Deep Venous Reflux	2.38	2.16	0-7
Is contraindicated by Deep Venous Obstruction	7.00	3.35	0-10
Should involve Direct injection of the IP	2.23	2.66	0-10
Should involve Injection into a communicating vessel above the fascia	8.65	2.06	0-10
Percentage of IPS closed completely with UGECA	75.88 %	18.87	30-100

Posters

In relation to UGECA technique the majority of respondents recommended use of foam sclerosant (Type = polidocanol or soltradecol; strength = 1-3%; Volume = 0.5 - 3 cc)

Conclusions: Based on the consensus of a group of expert phlebologists, the current opinion regarding incompetent perforator veins is that: 1) The presence of IPS is important in the pathogenesis of lipodermatosclerosis, venous ulcers, and varicose veins, 2) ablation of IPS may improve deep venous function, lipodermatosclerosis, ulcer healing, and edema, and reduce ulcer recurrence, and 3) UGECA using foam sclerosant into a communicating vessel above the fascia is the most commonly recommended technique for IP ablation.

P-28 Non-Saphenous Approach To Varicose Veins With Foam Sclerotherapy

V. Cheng - San Diego Vein Institute, Encinitas, CA

Background: Foam Sclerotherapy is emerging as effective treatment for varicose veins but ultrasound-guided access to the Saphenous Vein is the usual treatment approach. This study was done to see if vascular access through an available varix would simplify the procedure.

Methods: During the five-year period prior to June 2007, 1648 patients with venous disorders were evaluated and 408 patients had reflux in the great Saphenous vein. One hundred of the most recent patients (126 limbs) with primary varicose veins, CEAP class 2 with great Saphenous reflux were chosen for this analysis. There were 77 women (mean age 34 years) and 23 men (mean age 46 years). Exclusions were patients with venous leg ulcers, venous Angiomata, Klippel-Trenaunay syndrome, recurrent varicose veins and vascular malformations.

Vascular access was gained through an available varix and 1 to 3 needle sites were selected in each treated extremity. Ultrasound monitoring assured intravascular access for the 1-2% Polidocanol foam made by the Tessari technique. A Compounding Pharmacy prepared the Polidocanol for each patient and foam volumes were kept below 16 ml for each treated extremity. Effective 20-30 mm and 30-40 mm Hg stockings and supplemental focal compression with long stretch elastic bandaging for 48 to 72 hours excluded trapped blood and insured vein sealing. DVT surveillance at 7days and treatment evaluation at 7 and 28 days completed the procedure.

Results: Great Saphenous vein obliteration was achieved in 82% of the extremities, reversal of reflux in an additional 9%, and satisfactory correction of varicose veins in all extremities in an average of 2.8 treatment sessions. Adverse events were absent because of elevation of the legs 45° for 10 minutes following instillation of the foam. No iliofemoral or crural thromboses were detected by the surveillance. No patient required anesthesia or analgesia.

Conclusions: Satisfactory treatment of varicose veins can be achieved without using ultrasound access through the Great Saphenous vein. The procedure is considerably simplified and is less time consuming using direct visual vascular access through a varix rather than ultrasound guided Saphenous access.

P-29 Case Report: Epitheloid Hemangioendothelioma of the Common Femoral Vein

M. Lebow, A. Hurd, D. Cassada, M. Freeman, O. Grandas, S. Stevens, M. Goldman - University of Tennessee, Knoxville, TN

Background: Symptomatic unilateral leg swelling is a common complaint in patients with post-thrombotic syndrome. A 45 year old female with deep venous thrombosis 15 years earlier presented with left lower extremity swelling and pain. CT scan and venogram revealed a short-segment occlusion within the right common femoral vein and abundant collaterals.

Methods: A discrete solid mass occluding the vein was identified at operation. The mass was 1cm x 2cm, well circumscribed and adherent to the posterior wall of the vein. Because of its atypical appearance a complete oncologic vein resection was carried out with proximal and distal margins. A greater saphenous vein panel graft was used to reconstruct the femoral vein.

Results: Formal histologic analysis revealed Epithiloid Hemangioendothelioma (EH), completely excised. EH is a rare intravascular sarcoma with a predilection towards hematgenous and lymphatic metastasis and a significant mortality rate. Like other sarcomas it is resistant to chemotherapy and radiation. On clinical follow-up the leg swelling resolved and a metastatic work-up was negative. She will undergo lifelong surveillance for recurrence.

Conclusions: Primary vascular malignancies are an uncommon cause of venous occlusion. The ability to detect subtle differences in presentation coupled with a high index of suspicion helps in selecting patients for surgical excision and reconstruction for venous occlusion. When vasculogenic tumors are discovered, adherence to sound oncologic principals during excision minimizes the risk of recurrence and need for re-operation.

P-30 A Report of Two Rare Cases of Venous Aneurysms Involving the Lesser Saphenous Venous System

S. Chen, A. N. Bowser, W. D. Clouse, C. Johnson, T. E. Rasmussen - Wilford Hall Medical Center Lackland AFB, San Antonio, TX

Background: Venous aneurysms are rare and reported nearly exclusively in the deep venous system.

Methods: We present two cases of venous aneurysms involving the superficial venous system of the leg.

Results: A lesser saphenous vein aneurysm encountered in a 46 y/o female during evaluation of symptomatic leg varicosities. Examination revealed varicose veins of the leg (C2) and duplex demonstrated a 2.2cm lesser saphenous vein aneurysm with reflux. CT angiogram confirmed the aneurysm (Figure 1) and the patient underwent resection of the lesser saphenous vein aneurysm and phlebectomy of the varicose veins.

Case 2: A 40 y/o male with right leg varicosities and edema (C3) who had undergone ligation of the great saphenous vein with phlebectomy in the past. This patient had recurrence of right leg varicosities and persistent symptoms. Duplex revealed a large venous aneurysm in continuity with the popliteal and lesser saphenous veins. Duplex confirmed reflux in both the femoropopliteal and aneurysmal segment of lesser saphenous. The patient underwent surgical management of the aneurysm with phlebectomy.

Conclusions: These cases represent rare instances of venous aneurysms involving the superficial venous system of the leg. Specifically, lesser saphenous venous aneurysms have rarely been reported. While longstanding reflux and chronic venous hypertension may play a role in venous distension in general, these cases appear to represent distinct aneurysms with unique vessel morphology. Although venous aneurysms are rare they should be considered in the differential diagnosis of chronic venous insufficiency. The diagnosis and treatment of superficial venous aneurysms proceeds along the same lines as other superficial venous disorders.



P-31 Multimodal Endovascular - Open Surgical Approach To Phlegmasia Cerulea Dolens of the Upper Extremity: A Case Report

N. Patel, A. Puggioni, X. Li, A. Hingorani, A. Shiferson, V. Tran, E. Ascher - Maimonides Medical Center, Brooklyn, NY

Background: There are at least 250,000 recognized cases of DVT in the United States per year, of which approximately only 2-4% involve the upper extremity. Upper extremity DVT leading to phlegmasia cerulea dolens (PCD) occurs only in an estimated 2-5% of these cases, frequently leading to tissue loss and death. We report on a patient who developed deep venous thrombosis in the upper extremity resulting in PCD gangrene of the digits.

Methods: A 77 year old female was hospitalized following a severe respiratory tract infection. She was bed-ridden for many years due to poliomyelitis and a previous stroke. Shortly after admission patient sudden developed massive edema and cyanosis of her right upper extremity (Fig.1). Duplex ultrasound showed acute thrombosis of the brachial and cephalic veins as well as radial artery occlusion. Intravenous antibiotics, isotonic fluids, and heparin were administered. Emergency upper extremity phlebography via brachial vein cut down and rheolitic suction thrombectomy of the brachial and axillary veins was performed. The radial artery was explored, along with a forearm fasciotomy. A thrombolysis catheter was left in the right brachial vein for overnight tPA infusion (Fig.2); The limb was kept elevated.

Results: At 24 hours there was significant edema resolution and repeated venogram showed patent deep upper extremity veins. The upper extremity appearance improved over the following weeks (Fig.3) and regained acceptable motor function. Residual dry gangrene of fingertips was managed conservatively (Fig.4).

Conclusions: The use of a multimodality approach, surgical and interventional, was successful in treating the acute venous obstruction and relieving arterial compression, thereby preventing further fluid sequestration and limb loss.









P-32 Relation Between Number of Pregnancies and Great Saphenous Vein Diameters

S. X. Salles-Cunha¹, N. Morrison² - ¹CompuDiagnostics, Inc, Phoenix, AZ; ²Morrison Vein Institute, Phoenix, AZ

Background: Number of pregnancies has been mentioned as a risk factor for chronic venous insufficiency. We have related reflux to diameter of the great saphenous vein (GSV). This analysis investigated if number of pregnancies correlated with GSV diameter.

Method: Ultrasound evaluation of the GSV was performed during a voluntary service provided by American medical personnel in Guayaquil, Ecuador. Women who perceived they had leg venous problems were evaluated. Median number of pregnancies of 178 women, 51 ± 3 years of age, was 4 (range 0-15). Minimum number of women per subgroup of 0, 1, 2, $\frac{1}{4}$, 8, 9, >10 pregnancies was 5. GSV diameter in mm was measured at mid thigh with the patient standing. Statistical analysis included calculation of correlation coefficients and t-tests.

Results: Correlation coefficient between number of pregnancies and left GSV diameter calculated for the entire data set was low: 0.06. The correlation coefficient calculated for the average GSV diameter of each subgroup increased to 0.48. GSV diameter was smallest for women without pregnancies, $2.6\pm0.5 \text{ mm}$ (P<.001). GSV diameter of women having one pregnancy, $2.8\pm1.6 \text{ mm}$, was not significantly different than subgroups with less (P=.30) or more pregnancies (P=.17). Largest GSV diameters were: $4.2\pm3.4 \text{ mm}$ (N=9 pregnancies), $4.1\pm2.5 \text{ mm}$ (N>10), $4.1\pm2.8 \text{ mm}$ (N=2) and $4.0\pm2.8 \text{ mm}$ (N=5). The average GSV diameters for >9 or 1-8 pregnancies were not significantly different (P=.50).

Conclusions: GSV diameter increased with one pregnancy. Otherwise, the GSV diameter was not related to the number of pregnancies.

7:30 pm

THE FORUM FINALE

Awards, Dinner, Entertainment & More

NOTES

AMERICAN VENOUS FORUM

Alphabetical Roster

Honorary Members

Allegra, Claudio

S. Giovanni Hospital-Angiology Dept 26 Via Del Colosseo Rome, 00184 Italy Telephone: 39-0-6485527 Fax: 39-0-677055582 Email: allegra@mclink.lt

Bergqvist, David (Agneta)

University of Uppsala Academic Hospital Vasc. Surg. Uppsala, S-751 85 Sweden Telephone: 46-1-8664633 Fax: 46-1-8664632 Email: mona.bjorklund@kirurgi.uu.se

Bollinger, Alfred (Verena Elizabeth)

University of Zurich Trubelstr 31 Strafa, CH-8712 Switzerland Email: boll@goldnet.ch

Browse, Norman L (Jeanne)

Corbet House Butes Lane, Alderney Channel Islands, GY9 3UW United Kingdom Telephone: 44-1-481823716

Burnand, Kevin G

St Thomas Hosp, Academic Dept of Surgery 1st FIr North Wing, Lambeth Palace Road London, SE1 7EH United Kingdom Telephone: 44-2-076339405 Fax: 44-2-079288742 Email: kevin.burnand@kcl.ac.uk

Coleridge Smith, Philip D

Thames Valley Nuffield Hospital Wexham Street Wexham, SL3 6NH United Kingdom Telephone: 44-2-076368333 Fax: 44-2-076799413

Enrici, Ermenegildo A (Maine Moya)

Remedios de Escalada 2339 (1640) Martinez Bs.As Buenos Aires, 01123 Argentina Telephone: 54-1-147425440 Fax: 54-1-147425440 Email: enrici@colmed4.com.ar

Hirsh, Jack

Hamilton Civic Hosp Research Ctr 711 Concession St Hamilton, ON L8V 1C3 Canada Telephone: 90-5-5272299 Fax: 90-5-5752646 Email: ihirsch@thrombosis.hhscr.org

Hobbs, John T (Marianne)

4 Upper Wimpole St London, W1G 6LF United Kingdom Telephone: 20-7-3232830 Fax: 20-7-2242930 Email: john.t.hobbs@bropenworld.com

Natali, Jean P

17 rue Lamennais Paris, F-75008 France Telephone: 14-2-895439 Fax: 14-3-590100 Email:

Nicolaides, Andrew N (Lala)

Vascular Screening and Diagnostic Centre 2 Kyriacou matsi Street, Ayios Dhometios Nicosia, 01683 Cyprus Telephone: 35-7-22780543 Fax: 35-7-22780553 Email: anicolai@cytanet.com.cy

Partsch, Hugo

Medical University Baumeisterg 85 Vienna, A1160 Austria Telephone: 43-1-4855853 Fax: 43-1-4800304 Email: hugo.partsch@meduniwien.ac.at

Perrin, Michel

Clinique Du Grand Large 26 Chemin de Decines Chassieu, 69680 France Telephone: 33-4-72057266 Fax: 33-4-72057274 Email: m.perrin.chir.vasc@wanadoo.fr

Rabe, Eberhard

Klinik und Poliklinik fur Dermatologie Sigmund Freud Str. 25 Bonn, D-53105 Germany Telephone: 22-8-2875370 Fax: 22-8-2874333 Email: eberhard.rabe@ukb.uni-bonn.de

Ruckley, C. Vaughan

University of Edinburgh 1 Mayfield Terrace Edinburgh, EH9 1 RU United Kingdom Telephone: 13-1-6678678 Email: vaughan.ruckley@btinternet.com

Schmid-Schonbein, G. W.

University of California, San Diego 9500 Gilman Dr, Bioengr 0412 La Jolla, CA 92093-0412 USA Telephone: 61-9-5344272 Fax: 61-9-5345722

Thulesius, Olav

University Hospital Fac of Health Sciences Linkoping, S-581 85 Sweden Telephone: Fax: 46-1-3145949 Email: thulesius@juno.com

ACTIVE MEMBERS

Abai, Babak

UMDNJ-NJMS Division of Vascular Surgery 150 Bergen Street, E401 Newark, NJ 07101-1709 USA Telephone: 978-972-6295 Fax: 978-972-0092 Email: khoramdin@gmail.com

* Abbott, William M (Cynthia)

Massachusetts General Hospital 275 Charles St, Warren 901 Boston, MA 02114 USA Telephone: 617-726-8250 Fax: 617-726-3322 Email: wabbott@partners.org

AbuRahma, Ali F (Marion)

R C Byrd Health Sci Ctr of WVU 3110 MacCorkle Ave SE, Charleston, WV 25304 USA Telephone: 304-347-1306 Fax: 304-556-3823 Email: ali.aburahma@camc.org

Adelman, Mark A (Christie)

University Vascular Associates 530 1st Ave, #6F New York, NY 10016 USA Telephone: 212-263-7311 Fax: 212-263-7722 Email: mark.adelman@med.nyu.edu

Agarwal, Gautam

Mayo Clinic Gonda 4 South Vascular Surgery 200 1st St., SW Rochester, MN 55905 USA Email: gautam40@hotmail.com

Almeida, Jose Ignacio (Yvette)

Miami Vein Center 1501 South Miami Avenue Miami, FL 33129 USA Telephone: 305-854-1555 Fax: 305-854-1166 Email: jia@miamiveincenter.com

* Alpert, Joseph (Jane)

4 Top Gallant Cir. Savannah, GA 31411-2720 USA Telephone: 912-598-8287 Email: jalpert375@bellsouth.net

Anderson, Robert

Vein Centers for Excellence of Des Moines 1300 37th Street, Suite 3 West Des Moines, IA 50266 USA Telephone: 515-223-0592 Fax: 515-223-8316 Email: boba@veincenters.com

Angle, Niren

Univ. of California at San Diego 200 W. Arbor Drive San Diego, CA 92103 USA Telephone: 619-543-6980 Fax: 619-543-2615 Email: nangle@ucsd.edu

Araki, Clifford T (Linda)

St. Claire's Hospital 25 Pocono Rd. Denville, NJ 07834 USA Telephone: 973-625-6723

Arata, Michael

South Coast Vein Care 20162 Birch St., Suite 250 Newport Beach, CA 92660 USA Telephone: 949-706-3355 Fax: 949-209-2051 Email: admin@southcoastveincare.com

Arbid, Elias J (Rita)

Commonwealth Surgical Assoc. 3640 High Street Portsmouth, VA 23707 USA Telephone: 757-397-2383 Fax: 757-387-5201 Email: arbid@massmed.org

Ascher, Enrico (Katia)

Maimonides Med Ctr, Vasc Surgery 4802 Tenth Ave. Brooklyn, NY 11219 USA Telephone: 718-283-7957 Fax: 718-635-7050 Email: eascher@maimonidesmed.org

Baldwin, John C

The CBR Institute for Biomedical Research 200 Longwood Avenue Boston, MA 02115 USA Telephone: 617-278-3000 Fax: 617-278-3131 Email: baldwin@cbr.med.harvard.edu

Balkany, Louis

1614 So. Byrne Rd., Suite FF Toledo, OH 43614-3403 USA Telephone: 419-382-9425 Fax: 419-382-9427 Email: loubalkany@aol.com

Balshi, James D (Jill)

Progressive Physician Assoc, Inc. 3735 Nazareth Rd, #206 Easton, PA 18045 USA Telephone: 610-252-8281 Fax: 610-253-5321 Email: jbalshi@ppamail.com

* Barker, Wiley F (Nancy)

29129 Paiute Drive Agoura, CA 91301 USA Telephone: 818-865-9904 Fax: 818-865-9901 Email: wbarker@charter.net

* Baron, Howard C (Joan)

75 Central Park West 13D New York, NY 10023 USA Telephone: 212-362-0990

Bassiouny, Hisham S

University of Chicago 5841 So Maryland St, MC 5028 Chicago, IL 60637 USA Telephone: 773-702-6128 Fax: 773-702-0863 Email: hbassiou@surgery.bsd. uchicago.edu

Beavers, Frederick P (Cynthia Long)

Washington Hospital Center 110 Irving St. NW Washington, DC 20010 USA Email: suavejazz@hotmail.com

* Beebe, Hugh G (Carin Starr)

Dartmouth Hitchcock Medical Center One Medical Center Drive Lebanon, NH 03756 Telephone: 419-471-2088 Fax: 419-479-6980 Email: hbeebe@jvc.org

* Bergan, John J (Elisabeth)

9850 Genesee Ave, #410 La Jolla, CA 92037 USA Telephone: 858-550-0330 Fax: 858-550-0676 Email: jbergan@popmail.ucsd.edu

Bernhard, Victor M (Suzan)

3627 Grand Valley Canal Road Palisade, CO 81526 USA Telephone: 970-464-4653 Fax: 970-464-4654 Email: bernhard@surgery.bsd. uchicago.edu

Binnington, H. Bradley (Jeannine)

5032 Bischoff Ave. St Louis, MO 63110-3102 USA Telephone: 314-773-2830 Email: bbinnington@sbcglobal.net

Bjarnason, Haraldur (Katrin Frimannsdotter)

Mayo Clinic - Vascular and Interventional Radiology 200 First Street, SW Rochester, MN 55902 USA Telephone: 507-255-8454 Fax: 507-255-7872 Email: bjarnason.haraldur@mayo.edu

Blebea, John (Judy)

Temple University Hospital 3401 N Broad St, Parkinson Pav #433 Philadelphia, PA 19140 USA Telephone: 215-707-3622 Fax: 215-707-5901 Email: blebeaj@tuhs.temple.edu

* Blumenberg, Robert M (Gayle)

2259 Algonquin Rd Schenectady, NY 12309 USA Telephone: 518-393-7700

Bohannon, W. Todd

Scott & White Memorial Hospital & Clinic 2401 South 31st St. Temple, TX 76508 USA Telephone: 254-724-0657 Fax: 254-724-5978 Email: wbohannon@swmail.sw.org

* Boland, James P

RC Byrd Health Sciences Ctr 3110 MacCorkle Ave SE Charleston, WV 25304 USA Telephone: 304-347-1333

§ Bonawitz, Cara A

Medical Center Radiologists 6330 N. Center Dr, Bldg 13, Suite 220 Norfolk, VA 23502 USA Telephone: 757-466-0089 Fax: 757-466-8017 Email: cabonawitz@cox.net

Bradbury, Andrew W (Gillian)

University Department of Vascular Surgery Flat 5, Netherwood House Solihull Hospital Solihull B91 2JL United Kingdom Telephone: 44-1-214245086 Fax: 44-1-214245086 Email: andrew.bradbury@btinternet.com

Brown, O. William (Susan)

William Beaumont Hospital 31700 Telegraph Rd, #140 Bingham Farms, MI 48025 USA Telephone: 248-433-0881 Fax: 248-433-1628 Email: owbmd@aol.com

Brown, Kellie

Medical College of Wisconsin 9200 W. Wisconsin Ave MilwaUnited Kingdomee, WI 53226 USA Telephone: 414-805-9160 Email: krbrown@mcw.edu

Buchbinder, Dale (Sharon)

Greater Baltimore Med Ctr 6569 No Charles St, #701 Towson, MD 21204-6832 USA Telephone: 410-849-2393 Fax: 410-849-3435 Email: dbuchbin@gbmc.org

Buckman, Jeffrey (Myrna)

Vascular Diagnostics 1600 Dempster, #105 Park Ridge, IL 60068 USA Telephone: 847-298-7876 Fax: 847-298-7886 Email: j_buckman@msn.com

* Bulkin, Anatoly

SDIVA 488 E. Valley Pkwy, Suite 404 Escondido, CA 92025 USA Telephone: 760-739-7666 Fax: 760-739-7633 Email: abulkinmd@sdiva.com

§ Bush, Ruth

Michael E. DeBakey VA Medical Center 2002 Holcombe Blvd. Houston, TX 77030 USA Telephone: 713-794-7892 Fax: 713-794-7352 Email: rbush@swmail.org

Caggiati, Alberto (Sonia)

Department of Anatomy, University "La Sapienza" Via Borelli 50 Rome I-00153 Italy Email: alberto.caggiati@uniroma1.it

Calcagno, David (Elizabeth)

Calcagno and Rossi Vein Treatment Center 2025 Technology Parkway, Suite 304 Mechanicsburg, PA 17050 USA Telephone: 717-763-0510 Fax: 717-761-6081 Email: Vascularpc@msn.com

Calligaro, Keith D (Ina Lee)

Pennsylvania Hospital 700 Spruce St, #101 Philadelphia, PA 19106 USA Telephone: 215-829-5000 Fax: 215-627-0578 Email: kcalligaro@aol.com

Cambria, Robert A (Emily)

Medical College of Wisconsin 9200 W. Wisconsin Avenue Milwaukee, WI 53226 USA Telephone: 414-456-6970 Fax: 414-456-6216 Email: rcambria@mail.mcw.edu

* Cannon, Jack A (Helen)

25132 Via Pacifica Dana Point, CA 92629-2049 USA Telephone: 949-481-3328 Email: jac12@cox.net

Cantelmo, Nancy L (Michael Rauworth)

Massachusetts General Hospital One Hawthorne Place, Suite 111, H01 Boston, MA 02114 USA Telephone: 617-726-4464 Email: nlc31@comcast.net

* Caprini, Joseph A (Stella)

Evanston Northestern Healthcare 9977 Woods Drive Skokie, IL 60077 USA Telephone: 847-663-8050 Fax: 847-663-8054 Email: jcaprini2@aol.com

Carman, Teresa L

Cleveland Clinic Foundation 9500 Euclid Ave Desk S-60 Cleveland, OH 44195 USA Telephone: 216-445-5454 Fax: 216-444-7370 Email: tcarmanmd@aol.com

Carney, Wilfred I (Joan)

43 Acoaxet Rd. Westport, MA 02790 USA Telephone: 508-636-5405 Fax: 401-868-2303 Email: wilfredcarneymd@msn.com

Carr, Sandra C (Michael)

William S. Middleton Veterans Hos. 2500 Overlook Terrace, Ste B7054 Madison, WI 53705 USA Telephone: 608-263-1388 Fax: 608-280-7098 Email: carr@surgery.wisc.edu

Castronuovo, John J (Malin)

York Hospital, Surgery 1001 S. George St. York, PA 17405 USA Email: john.castronuovo@mmh.ahsys.org

Cazaubon, Nichele

Cabinet d'Angeiologie 48 rue St. Didier Paris 75116 France Telephone: 33-1-47271063 Fax: 33-1-47272147 Email: micazang@noos.fr

Cerveira, Joaquim J

Kaiser Permanente 13562 Cantara St, Surgery 201 Panorama City, CA 91402 USA Telephone: 818-375-3195 Email: jjc41@hotmail.com

Chang, Benjamin B (Heather)

The Vascular Group, PLLC 43 New Scotland Ave, MC 157 Albany, NY 12208 USA Telephone: 518-262-8720 Fax: 518-262-6720 Email: changb@albanyvascular.com

* Chang, John B (Lucy)

Long Island Vascular Center 1050 Northern Blvd Roslyn, NY 11576 USA Telephone: 516-484-3430 Fax: 516-484-3482 Email: jbchangmd@aol.com

Cheng, Van Le

Vein Institute of La Jolla 9850 Genesee Ave., Ste. 410 La Jolla, CA 92037 USA Telephone: 858-550-0330 Fax: 858-550-0676 Email: vanlecheng2003@yahoo.com

Cherry, Kenneth J (Robin)

University of VA Hospital PO Box 800679 Charlottesville, VA 22908 USA Telephone: 434-243-7052 Fax: 434-982-1026 Email: kjc5kh@virginia.edu

Cho, Jae-Sung (Michelle)

200 Lothrop St, PUH A1011 Pittsburgh, PA 15213 USA Telephone: 412-648-4000 Email: chojs@msx.upmc.edu

Clagett, G. Patrick (Nancy)

Univ of TX SW Medical Center 5323 Harry Hines Blvd. Dallas, TX 75390-9157 USA Telephone: 214-648-3516 Fax: 214-648-2790 Email: patrick.clagett@utsouthwestern.edu

Collins, David E.

Collins Vein & Laser Care PO Box 337 126 Trivette Dr. Pikeville, KY 41502 USA Telephone: 606-478-1407 Email: khli@tiusa.net

Comerota, Anthony J (Elsa)

Jobst Vascular Center 2109 Hughes Dr, #400-Conrad Jobst Twr Toledo, OH 43606 USA Telephone: 419-291-2088 Fax: 419-479-6980 Email: anthony.comerotamd@ promedica.org

Cordts, Paul R (Patricia Ann)

Office of the Surgeon General 5201 Brawner Place Alexandria, VA 22304-8645 USA Telephone: 703-681-0104 Fax: 703-681-6568 Email: paul.cordts@otsg.amedd.army.mil

Corson, John D (Tricia)

New Mexico VA Healthcare System 1501 San Pedro, SE Mail Drop 112 Albuquerque, NM 87108 USA Fax: 505-256-5743 Email: john.corson2@med.va.gov

Cranley, Robert D (Deborah)

Cranley Surgical Associates 3747 W Fork Rd Cincinnati, OH 45247-7548 USA Telephone: 513-961-4335 Fax: 513-961-4227 Email: taw@cranleysurgical.com

Criado, Enrique (Elena Camara)

University Hosp & Med Ctr HSC-18, 040 - Div Vascular Surgery Stony Brook, NY 11784-8191 USA Telephone: 631-444-2040 Fax: 631-444-8824 Email: ecriado@notes.cc.sunysb.edu

§ Daake, John W

The Reno Vein Clinic 1420 Holcomb Avenue, Suite A Reno, NV 89502 USA Telephone: 775-329-3100 Email: jdaake@renoveinclinic.com

Dalsing, Michael C (Rosa)

Indiana Univ. Med. School 1801 N. Senate Blvd. MPC II, #3500 Indianapolis, IN 46202 USA Telephone: 317-962-0280 Fax: 317-962-0289 Email: mdalsing@iupui.edu

Darling, R. Clement (Julie)

The Vascular Group, PLLC 43 New Scotland Ave, MC-157 Albany, NY 12208 USA Telephone: 518-262-8720 Fax: 518-262-6720 Email: darlingc@albanyvascular.com

Deak, Steven T (Kristen)

St Peter's University Hospital 37 Clyde Rd Ste #102 Somerset, NJ 08873-5034 USA Telephone: 732-873-0200 Fax: 732-873-0255 Email: sdeak@vascularnj.com

DeLaria, Giacomo A (Karen)

Scripps Clinic & Res Fnd 10666 Torrey Pines Rd La Jolla, CA 92037 USA Telephone: 858-554-8122 Fax: 858-554-6135 Email: delaria.giacomo@scrippshealth.org

* Delaurentis, Dominic A (Molly)

209 Sir Thomas Lunsford Drive Williamsburg, VA 23185 USA Telephone: 757-220-2592 Fax: 757-220-2987 * Denbo, Howard E (Lana) 45 Castro St., Ste. 138 San Francisco, CA 94114 USA Telephone: 415-776-9557 Fax: 415-922-0773 Email: hdenbo@sbcglobal.net

* Depalma, Ralph G (Maleva)

Dept. of Veterans Affairs 810 Vermont Ave NW, Rm 111B Washington, DC 20420 USA Telephone: 202-273-8505 Fax: 202-273-9108 Email: rgdepalma@mail.va.gov

* Deweese, James A (Patricia)

University of Rochester 601 Elmwood Ave Rochester, NY 14642 USA Telephone: 716-275-2721 Fax: 716-244-7171 Email: deweesepnj@aol.com

Dilling, Emery

Vein Solutions 6818 austin Center Blvd. Ste. 208 Austin, TX 78731 USA Telephone: 512-452-8346 Fax: 512-795-8346 Email: edilling@ctvstexas.com

§ Dion, Yves M (Marie)

Hopital St-Francois d'Assise 10 de l'Espinay Quebec QC G1L 3L5 Canada Email: dion.yves@videotron.ca

Donaldson, Magruder C (Jennifer)

Metro West Medical Center 85 Lincoln Street Framingham, MA 01702 USA Telephone: 508-383-1553 Fax: 508-383-1746 Email: m.donaldson@tenethealth.com

Donayre, Carlos E (Dorene)

Harbor/UCLA Medical Center 2324 Colt Road Rancho Palos Verdes, CA 90275 USA Telephone: 310-222-2704 Fax: 310-787-1889 Email: cdonayre@cox.net

* Dosick, Steven M (Sandra)

Veinsolutions, Toledo 2109 Hughes Dr, #550 Toledo, OH 43606-3856 USA Telephone: 419-291-2090 Fax: 419-479-6135 Email: smdosick@hotmail.com

* Duffy, David M

4201 Torrance Blvd, #710 Torrance, CA 90503-4511 USA Telephone: 310-370-5679 Fax: 310-214-2071 Email: info@drdavidmduffy.com

Durham, Joseph R (Marianne)

10347 So Longwood Drive Chicago, IL 60643 USA Telephone: 708-633-2800 Fax: 708-799-2261 Email: drhoser@aol.com

Edwards, James M (Michele Mass)

Portland VAMC (P-8-VS) 3710 US Veterans Hospital Rd Portland, OR 97207 USA Telephone: 503-220-8262 Fax: 503-220-3415 Email: edwardsj@ohsu.edu

* Eklof, Bo G (Monica)

University of Lund, Sweden Batteritorget 8 Helsingborg SE 252-70 Sweden Telephone: 464-226-0728 Email: moboek@telia.com

Eldrup-Jorgensen, Jens

The Maine Surgical Group 887 Congress St., Ste. 400 Portland, ME 04102 USA Telephone: 207-774-6368 Fax: 207-774-9388 Email: jensjorg@aol.com

Elias, Steven (Maria)

Englewood Hospital & Medical Center 350 Engle St. Englewood, NJ 07631 USA Telephone: 201-816-0666 Fax: 201-894-9951 Email: veininnovations@aol.com

* Elliott, Joseph P (Donna)

3282 Woodview Lake Rd West Bloomfield, MI 48323 USA

Elmore, Frederick A (Debra)

7131 No Eleventh St, #101 Fresno, CA 93710 USA Telephone: 559-435-0717 Fax: 559-435-9105 Email: jennifer@cvvein.com

Engle, Jennifer S (Paul S. Hartley)

3290 West Big Bear Road, Suite 410 Troy, MI 48084 USA

* Ernst, Calvin B (Elizabeth)

1 Greythorne Woods Circle Wayne, PA 19087 USA Telephone: 610-688-3445 Fax: 610-688-6690 Email: cbernst@earthlink.net

Feied, Craig F

Washington Hospital Center 110 Irving Street, NW Washington, DC 20010 USA Telephone: 202-877-7574 Fax: 202-965-0705 Email: cfeied@ncemi.org

§ Felty, Cindy

Mayo Clinic Medical Center 200 SW First St Rochester, MN 55905 USA Telephone: 507-266-9737 Fax: 507-266-1617 Email: felty.cindy@mayo.edu

Fernandez, Bernardo B (Rosa)

Cleveland Clinic Florida 2950 Cleveland Clinic Blvd Weston, FL 33331-3609 USA Telephone: 954-659-5230 Fax: 954-659-5292 Email: fernanb@ccf.org

Ferrier, Frank (Iris)

Ferrier Management & Consulting 3091 Farmington Drive Atlanta, GA 30339 USA Telephone: 404-943-1341 Fax: 404-943-1830 Email: fferrier@charter.net

* Ferris, Ernest J

Univ of AR for Med Sciences 4301 W Markham, Slot 556 Little Rock, AR 72205 USA Telephone: 501-686-5744 Fax: 501-686-6900 Email: ferrisernestj@uams.edu

Finkelmeier, William R (Terri)

Carmel Medical Center 13450 N. Meridian, Suite 160 Carmel, IN 46032 USA Telephone: 317-582-7676 Fax: 317-582-7099 Email: bfrailey@corvascmds.com

Fisher, Jay B ((Fran)

Progressive Physician Assoc, Inc. 3735 Nazareth Rd, #206 Easton, PA 18042 USA Telephone: 610-252-8281 Fax: 610-252-8614 Email: jfisher@ppamail.com

Flanigan, D. Preston (Beth)

St Joseph Hospital, Orange, CA 1140 W La Veta Ave, #850 Orange, CA 92868 USA Telephone: 714-560-4450 Fax: 714-560-4455 Email: knife@cox.net

Flinn, William R Univ of Maryland Medical Systems 22 So Greene St, #N4W66 Baltimore, MD 21201 USA Telephone: 410-328-5840 Fax: 410-328-0717 Email: wflinn@smail.umaryland.edu

Flynn, William F (Therese)

William F. Flynn Jr. MD PC 22 Mill St, #301 Arlington, MA 02476 USA Telephone: 781-643-6313 Fax: 781-643-6316 Email: wflynnjrmd@aol.com

Fodera, Maria Elena

New York Surgical Assoc. P.C. 2235 Clove Rd. Staten Island, NY 10305 USA Telephone: 718-815-8100 Fax: 718-815-8200 Email: mefodera@yahoo.com

* Fogarty, Thomas J (Rosalee)

3270 Alpine Rd Portola Valley, CA 94028 USA Telephone: 650-854-1822 Fax: 650-854-2778 Email: tjf@fogartybusiness.com

Forrestal, Mark (Deborah Foley)

Northwest Vein Care 1430 N. Arlington Hts. Road, Suite 206 Arlington Heights, IL 60004 USA Telephone: 847-259-8226 Email: nwveincare@hotmail.com

Franz, Randall (Dawn)

Central Ohio Vascular Services 285 E.State Street, Suite 260 Columbus, OH 43215 USA Telephone: 614-855-0862 Email: rfranz2@ohiohealth.com

* Fronek, Arnost (Kitty)

8461 Whale Watch Way La Jolla, CA 92037 USA Telephone: 619-534-4270 Fax: 619-534-1690 Email: afronek@vcsd.edu

Frusha, John D (Velarie)

Vascular Surgery Associates 8595 Picardy Ave., Ste. 320 Baton Rouge, LA 70809-3675 USA Telephone: 225-769-4493 Fax: 225-766-3144 Email: jfrusha@brvsa.com

Furey, Patricia C (Douglas Goumas)

Surgical Care Group, PC 4 Elliot Way, Suite 302 Manchester, NH 03103 USA Telephone: 603-627-1887 Email: drpfurey@msn.com

Gagne, Paul (Elizabeth)

New York University Medical Center 530 First Avenue 6F New York, NY 10016 USA Telephone: 212-263-7311 Fax: 212-263-7722 Email: paul.gagne@med.nyu.edu

Gale, Steven S (Katia)

Veinsolutions, Toledo 2109 Hughes Dr, #550 Toledo, OH 43606-3856 USA Telephone: 419-291-2090 Fax: 419-479-6135 Email: ssgale@jvc.org

Gardner, Glenn P (Lynn)

Univ. of Missouri Healthcare One Hospital Dr. Surgery, DC077.00 Columbia, MO 65212 USA Email: gardner_glenn@hotmail.com

* Gaspar, Max R (Lia)

1780 St John Road, #48-C Seal Beach, CA 90740 USA Telephone: 562-799-3318 Fax: 562-429-0807 Email: mgaspar@usc.edu

Gasparis, Antonios P (Theodora)

Stony Brook, Surgery HSC T-18 Rm 040 Stony Brook, NY 11794-8191 USA Telephone: 631-444-1279 Fax: 631-444-8824 Email: antonio.gasparis@stoneybrook.edu

Gillespie, David L (Mary)

13 Lakenheath Court Potomac, MD 20854 USA Telephone: 202-782-9928 Fax: 202-782-3198 Email: david.gillespie@na.amedd.army.mil

Ginzburg, Enrique (Barbara)

Univ of Miami, Dept of Surgery PO Box 016960, (D-40) Miami, FL 33101 USA Telephone: 305-585-7529 Fax: 305-585-3076 Email: eginzburg@miami.edu

Giordano, Joseph M (Orfa)

Geo Washington Univ Hosp 2150 Pennsylvania Ave, NW Washington, DC 20037 USA Telephone: 202-741-3225 Fax: 202-994-0567 Email: dbrothers@msa.gwu.edu

Gloviczki, Peter

Mayo Clinic 200 First St SW Rochester, MN 55905 USA Telephone: 507-284-4652 Fax: 507-266-7156 Email: gloviczki.peter@mayo.edu

Gocke, John (Marita)

LaGrange Vascular Center 5201 S Willow Spring Rd Suite 200 LaGrange, IL 60525 USA Telephone: 630-829-3835 Fax: 708-579-4986 Email: jegndmd@ameritech.net

Goldman, Mitchell H (Margy)

Univ of TN Grad Sch of Med, Surgery 1924 Alcoa Highway, Box U-11 Knoxville, TN 37920 USA Telephone: 865-544-9244 Fax: 865-544-6958 Email: mgoldman@mc.utmck.edu

* Gomes, Mario N (Belinda)

4701 Ogletown Stanton Rd, Ste #1204 Newark, DE 19713 USA Telephone: 302-623-4530 Fax: 302-623-4522 Email: mgomes@christiancare.org

* Goodson, Spencer F (Mary)

Methodist Hospital of Indiana 1801 North Senate Blvd. #755 Indianapolis, IN 46202 USA Telephone: 317-923-1787 Fax: 317-929-6259

Gradman, Wayne S

Beverly Hills Vein Center 235 South McCarty Drive Beverly Hills, CA 90212 USA Telephone: 310-550-9200 Fax: 310-277-5045 Email: wayne@gradman.com

Granke, Kenneth (Deborah)

Detroit VA Medical Center 7080 Colony Dr. West Bloomfield, MI 48323 USA Telephone: 734-740-0461 Fax: 313-576-1002 Email: kgranke@yahoo.com

Green, Richard M (Barbara)

Lenox Hill Hospital 130 East 77th St 13th Floor New York, NY 10021 USA Telephone: 212-434-3400 Fax: 212-434-3410 Email: rgreen@lenoxhill.net

* Greenfield, Lazar J (Sharon)

University of Michigan 1327 Jones Dr. #201 Ann Arbor, MI 48105 USA Telephone: 734-936-6398 Fax: 734-998-0173 Email: lazarg@umich.edu

Gruneiro, Laura A (Alex)

Geisinger Specialty Clinics 1000 East Mountain Blvd. Wilkesbarre, PA 18711 USA Telephone: 570-821-2340 Fax: 570-826-7904

* Gruss, Jorg D (Elisabeth)

Kurhessisches Diakonissenhaus Goethestrasse 85 Kassel D-34119 Germany Telephone: 56-1-1002314 Fax: 56-1-1002319

Gueldner, Terry L.

Wisconsin Vein Center 940 Maritime Dr. Manitowoc, WI 54220 USA Telephone: 920-686-7900 Fax: 920-686-7985 Email: dr@gueldnermd.com

Hakaim, Albert G

Mayo Clinic 4500 San Pablo Rd, Vascular Surgery Jacksonville, FL 32224 USA Telephone: 904-953-2077 Fax: 904-953-7368 Email: hakaim.albert@mayo.edu

Hallett, John W

Roper St. Francis Heart & Vascular Center 316 Calhoun Street Charleston, SC 29401 USA Telephone: 843-720-5665 Fax: 843-727-3370 Email: johnjeb.hallett@rsfh.com

Hammond, Sharon L (Sterling)

Colorado Cardiovascular Surgical Associates 6282 So Netherland Way Aurora, CO 80016-1326 USA Telephone: 303-388-6461 Email: shamo39@aol.com

Harris, E. John (Leslie)

Stanford Univ Medical Ctr 300 Pasteur Dr, H-3637, Vasc Stanford, CA 94305-5642 USA Telephone: 650-723-8648 Fax: 650-498-6044 Email: edjohn@stanford.edu

* Harris, Edmund J (Marilyn)

555 Laurel Ave, Ste #605 San Mateo, CA 94401-4153 USA Telephone: 650-348-1414 Fax: 650-348-1414

Harris, Linda M (Norm Moser)

Millard Fillmore Hospital 3 Gates Circle, Dept of Surgery Buffalo, NY 14209 USA Telephone: 716-887-4807 Fax: 716-887-4220 Email: Imharris@acsu.buffalo.edu

Hasaniya, Nahidh W

Loma Linda University Medical Center 11175 Campus Street Suite 21121 Loma Linda, CA 92354 USA Telephone: 909-558-4354 Fax: 909-558-0348 Email: nahidh@pol.net

Haser, Paul B

UMDNJ - RWJMS, Vascular Surgery 1 Robert Wood Johnson Pl. MEB-54 New Brunswick, NJ 08903-0019 USA Telephone: 732-235-7816 Fax: 732-235-8538 Email: haserph@umdnj.edu

Henke, Peter K (Barbara)

Univ of MI Health System 1500 E Med Ctr Dr, 2210D Taubman Ctr Ann Arbor, MI 48109-0329 USA Telephone: 734-763-0250 Fax: 734-647-9867 Email: henke@umich.edu

Hill, Douglas

The Vein Treatment Centre 2004 14th Street NW, #207 Calgary, AB T2M3N3 Canada Telephone: 403-220-9353 Fax: 403-210-0593 Email: douglashill@shaw.ca

Hingorani, Anil P (Renu)

Maimonides Medical Center 4802 10th Ave, Admin Bldg Brooklyn, NY 11219 USA Telephone: 718-283-7957 Fax: 718-635-7050 Email: ahingorani@maimonidesmed.org

* Hobson, Robert W (Joan)

UMDNJ-NJ Medical School 30 Bergen St, ADMC Bldg 6, Rm 620 Newark, NJ 07107 USA Telephone: 973-972-6633 Fax: 973-972-5924 Email: hobsonrw@umdnj.edu

Hollier, Larry H (Diana)

LSU School of Medicine 533 Bolivar St. New Orleans, LA 70012 USA Telephone: 504-568-4009 Fax: 504-568-4008 Email: Ihholl@lsuhsc.edu

Hunter, Glenn C

University of Texas Medical Branc 301 University, 6.136 McCullough Galveston, TX 77555-0544 USA Telephone: 409-772-6366 Fax: 409-747-0966 Email: gchunter@utmb.edu

Hutto, John D.

University of Cincinnati 231 Albert Sabin Way Box 670558 Cincinnati, OH 45209 USA Telephone: 513-558-5367 Email: jd_hutto@yahoo.com

lafrati, Mark D (Jane Freedman)

New England Medical Center 750 Washington St, NEMC 1035 Boston, MA 02111 USA Telephone: 617-636-8094 Fax: 617-636-8003 Email: miafrati@tufts-nemc.org

Illig, Karl A (Juliet)

Univ of Rochester Med Ctr 601 Elmwood Ave, Box 652 Rochester, NY 14642 USA Telephone: 716-275-6772 Fax: 716-273-1077 Email: karl_illig@urmc.rochester.edu

Isaacs, Mark

Walnut Creek 1981 N. Broadway, Suite 427 Walnut Creek, CA 94596 USA Telephone: 925-945-8656 Fax: 925-945-8818 Email: misaacs@veinspec.com

Jamil, Zafar (Shireen)

St Michael's Medical Center 306 Dr M L King Jr Blvd, MS-45 Newark, NJ 07102 USA Telephone: 973-877-5059 Fax: 973-877-2954

Jarrett, Fredric (Esther)

UPMC-Shadyside 5200 Centre Ave, #716 Pittsburgh, PA 15232-1300 USA Telephone: 412-681-8720 Fax: 412-681-8713 Email: jarrettf@msx.upmc.edu

Johnston, Robert H (Sara)

Vein Clinics of Texas P.O. Box 3353 Victoria, TX 77903 USA Telephone: 361-570-8346 Fax: 512-582-5780 Email: bobyjohn@aol.com

Kabnick, Lowell S

New York University Medical School NYU Vein Center - 530 1st Ave. Suite 6D New York, NY 10016 USA Telephone: 212-263-8346 Email: doctlc@aol.com

Kalra, Manju

Mayo Clinic 200 First Street, SW Rochester, MN 55905 USA Telephone: 507-284-4494 Fax: 507-266-7156 Email: kalra.manju@mayo.edu

Kang, Steven S (Sylvia)

Reiss & Kang, M.D.,P.A. 9075 SW 87th Avenue, #414 Miami, FL 33176 USA Telephone: 305-598-0888 Fax: 305-598-3101 Email: vascular@bellsouth.net

Kanter, Alan

Vein Center of Orange County 250 East Yale Loop, Suite D Irvine, CA 92604-4697 USA Telephone: 949-551-8855 Fax: 949-527-8860 Email: veindoc@fea.net

Kasirajan, Karthikeshwar

Emory University Hospital 1364 Clifton Road NE, STE H-122A Atlanta, GA 30322 USA Telephone: 404-727-8407 Fax: 404-727-3316 Email: karthik_kasirajan@ emoryhealthcare.org

§ Kaufman, Steven L

Total Vein Care 1136 E. Stuart Street Suite 4102 Fort Collins, CO 80525 USA Telephone: 970-498-8346 Fax: 970-419-8346 Email: info@totalvein.net

Kazmers, Andris (Irene)

Petoskey Surgeons 560 W. Mitchell, Ste #140 Petoskey, MI 49770 USA Telephone: 231-487-1900 Fax: 231-487-2707 Email: akazmers@excite.com

* Kempczinski, Richard

3435 Golden Ave, Apt 201 Cincinnati, OH 45226 USA Telephone: 513-321-4724 Fax: 513-321-7350 Email: kemprf@fuse.net

Kent, K. Craig (Gina)

NY Presbyterian Hospital 525 East 68th St, Rm P-707, Box 197 New York, NY 10021 USA Telephone: 212-746-5192 Fax: 212-746-5812 Email: kckent@mail.med.cornell.edu

Kerr, Thomas M (Patricia)

4600 No Habana Ave, #28 Tampa, FL 33614 USA Telephone: 813-348-9088 Fax: 813-348-9310 Email: vasculardoc1@aol.com

* Kerstein, Morris D (Margaret)

1601 Kirkwood Highway Wilmington, DE 19805 USA Telephone: 610-527-4316 Fax: 610-520-9293

Killewich, Lois A

Univ of TX Med. Branch 301 University Blvd., Dept. of Surg. Room 6. 136 McCullough Galveston, TX 77555-0735 USA Telephone: 409-772-6366 Fax: 409-747-0966 Email: lakillew@utmb.edu

* Kistner, Robert L (Adelaide)

Beretania Medical Plaza 848 So. Beretania Street Suite 307 Honolulu, HI 96813 USA Telephone: 808-532-8346 Fax: 808-532-2240 Email: rlk@aloha.com

* Kloecker, Richard J (Phyllis)

8 Outer Ladue Drive St Louis, MO 63131 USA Telephone: 314-692-9100 Fax: 314-569-3119

Knott, Andrew W

Mayo Clinic 200 First Street SW 5-Joseph Rochester, MN 55905 USA Telephone: 507-255-7838 Email: knott.andrew@mayo.edu

* Konigsberg, Stephen F (Rhoda)

Highland Park Surgical Associates 31 River Rd Highland Park, NJ 08904 USA Telephone: 732-846-9500 Fax: 732-846-3931

Kritpracha, Boonprasit (Charuwan)

2109 Hughes Dr, Ste #400 Toledo, OH 43606 USA Telephone: 419-291-2080 Fax: 419-479-6980 Email: bkritpracha@jvc.org

Kurtoglu, Mehmet H.

Istanbul Medical Facility Emergency Surgery Capa, Topkapi Istanbul 34390 Turkey Telephone: 90-2-125742959 Email: metlevkurt@superonline.com

Labropoulos, Nicos

UMDNJ 150 Bergen Street, Room D-447 Newark, NJ 07101-1709 USA Telephone: 973-972-4138 Fax: 973-972-0433 Email: nlabrop@yahoo.com

Lal, Brajesh K (Priti)

UMDNJ, Div Vascular Surgery 185 S. Orange Ave., MSB-H570 Newark, NJ 07103 USA Telephone: 973-972-3736 Fax: 973-972-7425 Email: lalbk@umdnj.edu

Lalka, Stephen G (Valerie)

IN Univ Med Ctr, Wishard 1001 W 10th St, OPE 310A Indianapolis, IN 46202 USA Telephone: 317-962-0281 Fax: 317-962-0289 Email: sglalka@iupui.edu

Lall, Purandath

Mayo Clinic Dept. of Vascular Surgery 200 First St., SW Rochester, MN 55905 USA Telephone: 507-284-2511 Fax: 507-284-0161 Email: purandathlall@hotmail.com

* Lamesch, Alfred J

Clinic Dr Bohler 30 Rue de Luxembourg Goetzingen L-8360 Luxembourg Telephone: 44-9-06455

Laredo, James

Georgetown Univ. Hospital Dept. of Surgery 3800 Rservoir Rd, NW, 4 PHC Washington, DC 20007 USA Telephone: 202-444-2255 Fax: 202-444-6498 Email: jl393@georgetown.edu

Lauber, Andre F

Venenpraxis Unter Oter Egg No Lucerne, MD 06004 Switzerland Telephone: 41-4-13705570 Fax: 41-4-13705370 Email: lauber@venen-praxis.ch

* Lee, Byung-Boong (Hikyung)

Georgetown University 1830 Town Center Drive Suite 401 Reston, VA 20190 USA Telephone: 703-880-9500 Fax: 703-880-9598 Email: bblee38@comcast.net

Lemmon, Gary W (Kim)

Good Samaritan Hospital 2200 Philadelphia Drive, Med Education Dayton, OH 45406 USA Telephone: 937-278-6251 Fax: 937-276-8253 Email: gary.lemmon@wright.edu

Leon, Luis (Christine Renee Poock)

Loyola University Medical Center 2160 S. First Avenue Maywood, IL 60153 USA Telephone: 708-327-2236 Fax: 708-216-6300 Email: Ileon@lumc.edu

Liasis, Nikolaos E

Mesogion 109-111 Athens 11526 Greece Telephone: 30-2-106911668 Fax: 30-2-106911676 Email: nikos.liasis@euromedic.gr

Lin, Peter (Karla)

Baylor College of Medicine HVAMC-112 2002 Holcombe Blvd. Houston, TX 77030 USA Telephone: 713-794-7892 Fax: 713-794-7352 Email: plin@bcm.tmc.edu

* Lofgren, Eric P (Dorothy)

Mayo Clinic 200 First St SW Rochester, MN 55901 USA Telephone: 507-284-2511

* Lofgren, Karl A (Jean)

211 2nd St NW, Apt #1916 Rochester, MN 55901 USA Telephone: 507-284-2691

Lohr, Joann M (Michael Reardon)

Lohr Surgical Specialists 6350 Glenway Ave, #208 Cincinnati, OH 45211-6378 USA Telephone: 513-451-7400 Fax: 513-451-7888 Email: jlohr@lohrss.com

Long, John B (Teresa)

California Pacific Medical Center 3838 California St San Francisco, CA 94118 USA Telephone: 415-221-7056 Fax: 415-221-3583 Email: drilong@aol.com

Lumsden, Alan B (Terry Rice)

The Methodist Hospital Cardiovascular Surgery Dept. 6560 Fannin Street, Suite 1006 Houston, TX 77030 USA Telephone: 713-798-8412 Fax: 713-798-8632 Email: alumsden@tmhs.org

Lurie, Fedor (Galina)

Kistner Vein Clinic 848 South Beretania Street, Suite 307 Honolulu, HI 96813 USA Telephone: 808-532-8346 Fax: 808-532-2240 Email: flurie@kistnerveinclinic.com

Lynch, Thomas G (Jane)

Univ of NE Medical Center 9721 Spring St. Omaha, NE 68124 USA Telephone: 402-391-5811 Fax: 402-559-6749 Email: tomlynch@cox.net

Lynn, Richard A (Margrit Bessenroth-Lynn)

1411 No Flagler Dr, #9700 West Palm Beach, FL 33401-3413 USA Telephone: 561-655-1877 Fax: 561-655-6404 Email: rich549bux@aol.com

Maharaj, Dale A

12 Park View - Trincity Trinidad, West Indies Telephone: 868-640-4619 Fax: 518-262-6720 Email: dalemaharaj@hotmail.com

Mansour, M. Ashraf (Julie)

Michigan State University 4069 Lake Drive S.E., Suite 312 Grand Rapids, MI 49546-8816 USA Telephone: 616-459-8700 Fax: 616-459-0247 Email: ashmans2@aol.com

Marston, William A (Laurie)

3023 Burnett-Womack Building Department of Surgery University of North Carolina Chapel Hill, NC 27599-7212 USA Telephone: 919-966-3391 Fax: 919-966-2898 Email: sky@med.unc.edu

* Martin, Alfred J (Thomasine Alicia)

PO Box 4697 Santa Fe, NM 87502 USA Telephone: 505-82-01544 Fax: 505-982-0382 Email: ajmartinjr@msn.com

Martinez, Jeffrey M.

Peripheral Vascular Associates 111 Dallas St., Suite 200A San Antonio, TX 78205 USA Telephone: 210-225-7508 Fax: 210-225-1486 Email: jmartinez@pvasctx.com

Masuda, Elna M (Kevin Lui)

Straub Clinic & Hospital 888 So King St, Palma 5 Honolulu, HI 96813 USA Telephone: 808-522-4469 Fax: 808-522-4523 Email: elna.masuda@straub.net

Matsumura, Jon S (Amy)

NMFF 201 E. Huron St, Ste 10-105 Chicago, IL 60611 USA Telephone: 312-695-4857 Fax: 312-695-4955 Email: j-matsumura@northwestern.edu

Mattos, Mark A

Harper Hospital / Detroit Medical Center Vascular Surgery 3990 John Road Detroit, MI 48201 USA Telephone: 313-745-8637 Fax: 313-993-0244 Email: mmattos@dmc.org

McCarthy, Walter J (Mary)

Rush Presbyterian-St LUnited Kingdome's Hosp 1725 W Harrison, Rm 1156 Chicago, IL 60612 USA Telephone: 312-563-2762 Fax: 312-829-8680 Email: wmccart1@rush.edu

* Mckittrick, James E (Mehle)

649 Camino Campana Santa Barbara, CA 93111-1424 USA Telephone: 805-967-3282 Fax: 209-315-5808 Email: jmckinsb@aol.com

McLafferty, Robert B (Erica)

SIU Medical Center 800 N. Rutledge Street Suite D346 Springfield, IL 62702 USA Telephone: 217-545-7983 Fax: 217-545-2563 Email: rmclafferty@siumed.edu

Meissner, Mark H (Nancy)

University of Washington Medical Center Dept. of Surgery, Box 356410 1959 NE Pacific St., Room BB487 Seattle, WA 98915-6410 USA Telephone: 206-221-7047 Fax: 206-616-7495 Email: meissner@u.washington.edu

Menzoian, James O (Deborah Syah)

University of CT Health Center 263 Farmington, Ave. Farmington, CT 06030 USA Telephone: 860-679-7650 Fax: 860-679-4948 Email: jmenzoian@uchc.edu

Merchant, Robert F (Stephanie)

The Reno Vein Clinic 1420 Holcomb Ave, #A Reno, NV 89502-2960 USA Telephone: 775-329-3100 Fax: 775-329-3199 Email: doc@renoveinclinic.com

§ Meretei, Attila

Clinasys LLC 6797 Willow Wood Drive, #6036 Boca Raton, FL 33434 USA Telephone: 561-488-0422 Fax: 561-558-1358 Email: attila@clinasys.com

Merli, Geno J (Charlotte)

Jefferson Medical College 833 Chestnut St, Ste #701 Philadelphia, PA 19107 USA Telephone: 215-503-1022 Fax: 215-923-9239 Email: geno.merli@jefferson.edu

Messina, Louis M (Catherine)

Division of Vascular Surgery 55 Lake Avenue North Worcester, MA 01655-0337 USA Telephone: 508-856-5599 Email: moorer@surgery.ucf.edu

Mewissen, Mark W

St Luke's Vascular Center 2801 W Kinnickinnic Rvr Pkwy, #540 Milwaukee, WI 53215-3606 USA Telephone: 414-649-3599 Fax: 414-649-8140

Min, Robert J (Seri Ann Saltzman)

Weill Cornell Medical College 525 East 68th St. Room Starr 8a-37 New York, NY 10021 USA Telephone: 212-746-2520 Email: rjm2002@med.cornell.edu

Mintz, Bruce (Barbara Girz)

St Clare's Riverside Med Ctr 16 Pocono Rd, #313 Denville, NJ 07834 USA Telephone: 973-625-0112 Fax: 973-625-0721

Miskin, Barry M (Rita)

1926 Lenmore Dr. Palm Beach Gardens, FL 33410 USA Telephone: 561-745-7789 Fax: 561-745-4470 Email: miskinmd@aol.com

Monahan, Daniel L (Lynette Sue Monahan)

Vein Surgery & Treatment Center of No. California 1211 Pleasant Grove Blvd., Suite 120 Roseville, CA 95678-6971 USA Telephone: 916-791-8346 Fax: 916-791-8833 Email: danlmonahan@hotmail.com

Monedero, Javier Leal

Hospital Ruber Internacional C/ LA Maso N. 38 Madrid, AL 28034 Spain Telephone: 34-9-13875157 Fax: 34-9-13875158 Email: angiovascularlyz@ ruberinternacional.es

Moneta, Gregory L (Tracey)

OR Health Sciences Univ, Vasc 3181 SW Sam Jackson Pk Rd Portland, OR 97201-3098 USA Telephone: 503-494-7593 Fax: 503-494-4324 Email: monetag@ohsu.edu

Morasch, Mark D

Northwestern University Med School 201 E Huron St, #10-105, Vasc Surgery Chicago, IL 60611 USA Telephone: 312-695-2716 Fax: 312-695-4955 Email: mmorasch@nmh.org

Moritz, Mark W

Vein Institute of New Jersey 95 Madison Avenue Morristown, NJ 07960 USA Telephone: 973-539-6900 Fax: 973-588-4115 Email: mmoritz@vinj.us

Morrison, Nick (Terri)

Morrison Vein Institute 8575 E. Princess Dr., Suite 223 Scottsdale, AZ 85255 USA Telephone: 480-860-6455 Fax: 480-860-6679 Email: nickmorrison2002@yahoo.com

Muck, Patrick E (Sherry)

Good Samartian Hospital 375 Dixmyth Ave, 3rd FI, Surgery Cincinnati, OH 45220 USA Telephone: 513-232-8181 Fax: 513-624-2964 Email: pmuck@fuse.net

* Mulcare, Robert (Betsy)

9 Cedarwood Drive Greenwich, CT 06830 USA Email: drrjmulc@aol.com

Murray, James D

Kaiser Permanente - Vasc. Surg. 1011 Baldwin Pk. Blvd. Baldwin Park, CA 91706 USA Telephone: 626-851-6878 Fax: 626-851-6802 Email: james.d.murray@kp.org

Mutyala, Manikyam

374 Stockholm St. Brooklyn, NY 11237 USA Telephone: 718-486-4159 Email: mutyala68@hotmail.com

Myers, Jr., Daniel

University of Michigan 1150 W. Medical Center Drive MSRB II A570D Ann Arbor, MI 48109-0654 USA Email: ddmyers@umich.edu

§ Navarro, Felipe

North Ohio Heart Center 29325 Health Campus Drive, Suite 3 Westlake, OH 44145 USA Telephone: 440-617-2700 Fax: 440-808-8480 Email: fngvarro@nohc.com

Nazzal, Munier M.S. (Iman Mohamed)

Medical College of Ohio, Surgery 3064 Arlington Ave. Toledo, OH 43614 USA Telephone: 419-383-6810 Email: mnazzal@meduohio.edu

Neglen, Peter (Pamela)

River Oaks Hospital 1020 River Oaks Drive, Suite 480 Flowood, MS 39232 USA Telephone: 601-664-6680 Fax: 601-664-6694 Email: neglenmd@earthlink.net

Nicholls, Stephen (Elena Robinson)

Southwest Washington Thoracic and Vascular Surgery 200 NE Mother Joseph Place, Suite 300 Vancouver, WA 98664 USA Telephone: 360-514-1854 Email: snicholl@swmedicalcenter.com

Noel, Audra A

Mayo Clinic 200 First St SW, Gonda 4South Rochester, MN 55905 USA Telephone: 507-284-4751 Fax: 507-266-7156 Email: noel.audra@mayo.edu

Noppeney, Thomas (Jeanette)

Klinik Hallerwiese, Dept. of Surgery / Praxis fuer Gefaessmedizin Obere Turnstrasse 8-10 Nuremberg D-90429 Germany Telephone: 49-9-112706170 Fax: 49-9-112706181 Email: thoppeney.nbg@t-online.de

Nypaver, Timothy J (Michele)

Henry Ford Hospital 2799 W Grand Blvd, Vascular Surgery Detroit, MI 48202 USA Telephone: 313-916-3153 Fax: 313-916-3023 Email: tnypave1@hfhs.org

O'Byrne, Margaret G

10666 N. Torrey Pine Road SW 208 La Jolla, CA 92037 USA Telephone: 619-218-8980 Fax: 858-793-1632 Email: mastrodimos@aol.com

Oderich, Gustavo (Thanila Macedo)

Mayo Clinic 200 First Street SW Rochester, MN 55901 USA Email: oderich.gustavo@mayo.edu

O'Donnell, Thomas F (Carolyn)

New England Medical Center 750 Washington St, Box 259 Boston, MA 02111 USA Telephone: 617-636-5660 Fax: 617-636-5936 Email: todonnell@tufts-nemc.org

Olin, Jeffrey W (Joanie)

Mt Sinai School of Medicine One Gustave Levy PI, Box 1033 New York, NY 10029-6574 USA Telephone: 212-241-9454 Fax: 212-241-5107 Email: jeffrey.olin@msnyuhealth.org

Oliver, Mark A (Elise)

Morristown Memorial Hospital 182 South Street Morristown, NJ 07960 USA Telephone: 973-538-0165 Fax: 973-538-9344 Email: cdoppler@aol.com

O'Shea, Susan I (John)

DUnited Kingdome University Medical Center Erwin Rd, Box 3422 Durham, NC 27710 USA Telephone: 919-684-5350 Fax: 919-681-6160 Email: oshea005@mc.dUnited Kingdome.edu

Owens, Lewis (Kelly Anne)

CRL Surgical Associates 1490 Pantops Mountain Place Suite 100 Charlottesville, VA 22911 USA Telephone: 434-244-4580 Fax: 434-244-4579 Email: lewis.owens@mjh.org

Padberg, Frank T (Sharon)

Doctors Office Center 90 Bergen St., Ste. 2300 Newark, NJ 07103 USA Telephone: 973-676-1000 Fax: 973-395-7193 Email: padbergjr@aol.com

Paladugu, Ramesh

Plains Regional Medical Center 2200 Twenty First St. Clovis, NM 88101 USA Telephone: 505-769-6440 Fax: 505-769-6442 Email: rameshpal@pol.net

Pappas, Peter J (Nadine)

UMDNJ - University Hospital, Vascular Surgery 90 Bergen Street, Suite 7600 Newark, NJ 07101 USA Telephone: 973-972-6295 Fax: 973-972-0092 Email: pappaspj@umdnj.edu

* Paramo-Diaz, Marcelo

Av Alfonso Reyes 161 Mexico, DF 06140 Mexico Telephone: 525-515-3201 Fax: 525-516-5362

Pascarella, Luigi

University of California San Diego 9500 Gilman Dr. Bioengineering 0412 LaJolla, CA 92093-0412 USA Telephone: 858-538-2714 Fax: 858-550-0676 Email: pluigi@be-research.ucsd.edu

Passman, Marc A (Cora)

University of Alabama at Birmingham Section of Vascular Surgery BDB 503 1808 7th Avenue South Birmingham, AL 35294-0012 USA Telephone: 205-934-2003 Fax: 205-934-0024 Email: marc.passman@ccc.uab.edu

Patterson, Robert B

Providence Surgical Care Group 486 Silver Spring Street Providence, RI 02904 USA Telephone: 401-454-0690 Fax: 401-454-4281 Email: robert_patterson@brown.edu

Pavcnik, Dusan

Dotter Interventional Inst.,OHSU L342 630 SW Gaines Street Portland, OR 97239-3098 USA Telephone: 503-494-3669 Fax: 503-494-4258 Email: pavcnikd@ohsu.edu

Pearce, William H (Ann)

Northwestern Medical Faculty Fdn 201 East Huron #10-105, Vasc Surgery Chicago, IL 60611 USA Telephone: 312-926-7775 Fax: 312-695-4955 Email: wpearce@nmh.org

Peden, Eric

Baylor College of Medicine 1709 Dryden Street Suite 1500 Houston, TX 77030 USA Telephone: 713-798-8412 Fax: 713-798-8632 Email: epeden@bcm.edu

* Persson, Alfred V (May)

5 Dean Road Wellesley, MA 02481 USA Telephone: 781-235-6910 Fax: 781-4311632

* Pfeifer, John R (Jeanne)

University of Michigan, Venous Disease 19900 Haggerty Rd., #105 Livonia, MI 48152 USA Telephone: 734-432-7662 Fax: 734-432-7637 Email: pfeiferi@umich.edu

Phifer, Travis J

LSU Med Ctr, Dept Surgery 1501 Kings Hwy, PO Box 33932 Shreveport, LA 71130-3932 USA Telephone: 318-675-7770 Fax: 318-675-6141 Email: tphife@lsuhsc.edu

Pittaluga, Paul

Riveriera Vein Institute 10 Av. De Villeneuve Cagnes sur ner 06800 France Telephone: 33-4-92133413 Fax: 33-4-93207202 Email: paulpittaluga@hotmail.com

Polak, Joseph F

New England Medical Center 750 Washington St., Radiology Boston, MA 02111 USA Telephone: 617-636-6090 Fax: 617-636-0041 Email: jpolak@tufts-nemd.org

Pounds, Lori C (Kevin)

Univ of TX Medical Branch 301 Univ Blvd, 6.110 John Sealy Annex Galveston, TX 77555-0541 USA Telephone: 409-772-6369 Fax: 409-747-0966 Email: lori.pounds@utmb.edu

Powell, C. Steven (Melissa)

East Carolina Univ Schl of Med Dept of Surgery Greenville, NC 27858 USA Telephone: 252-816-4668 Fax: 252-816-3794 Email: powellc@mail.ecu.edu

Pringle, Timothy C

Good Samaritan Hospital 375 Dixmyth Ave, Hatton Rsrch 11J Cincinnati, OH 45220-2489 USA Telephone: 513-872-2785 Fax: 513-872-1549 Email: timothycp00@yahoo.com

Procter, Charles D (Elizabeth)

Surgical Specialists of Georgia 1250 Jesse Jewel Pkwy, #300 Gainesville, GA 30501 USA Telephone: 770-534-0110 Fax: 770-531-2423 Email: cdprocter@gmail.com

§ Proctor, Mary C (William)

Orthofix 1720 Bray Central Drive McKinney, TX 75069 USA Telephone: 214-578-2234 Email: maryproctor@orthofix.com

Proebstle, Thomas

Private Practice Zinkenbergweg 2 Hirschberg D-69493 Germany Telephone: 49-6-712065419 Fax: 49-6-201879660 Email: thomas.proebstle@web.de

Puggioni, Alessandra

Maimonides Medical Center 4703 8th Avenue Apartment 1 Brooklyn, NY 11220-1524 USA Telephone: 507-255-5123 Email: alpuggions2000@yahoo.com

Raffetto, Joseph D (Tamara)

VA Boston Healthcare System 1400 VFW Prkwy, Surgery 112, Vasc West Roxbury, MA 02132 USA Telephone: 857-203-5572 Fax: 857-203-5567 Email: joseph.raffetto@med.va.gov

Rai, Dinker B (Shakila)

555 Prospect Place Brooklyn, NY 11238 USA Telephone: 718-499-0202 Fax: 516-248-1547 Email: dbrai@aol.com

* Raju, Seshadri (Sybil)

Seshadri Raju, MD, PA 1020 River Oaks Drive, #420 Flowood, MS 39232 USA Telephone: 601-939-4230 Fax: 601-939-5210 Email: rajumd@earthlink.net

Ramnauth, Subhash C

Riverside Medical 401 Market Street, Suite 200 Steubenville, OH 43952 USA Telephone: 740-282-5000 Fax: 740-282-5233 Email: sram@riversidemds.com

* Ratliff, Jack L (Brenda)

410 University Pkwy, #2310 Aiken, SC 29801 USA Telephone: 803-648-1318 Fax: 803-642-7803

Razvi, Syed A (Tahera)

Caritas St. Elizabeth's Medical Center Medical Office Building 11 Nevins St, #308 Brighton, MA 02135 USA Telephone: 617-254-4200 Fax: 617-254-4242 Email: syed.a.razvi@verizon.net

Reed, Amy B

University of Cincinnati Div. of Vasc. Surgery 231 Albert Sabin Way Cincinnati, OH 45267 USA Telephone: 513-558-5367 Email: amy.reed@uc.edu

Rhodes, Jeffrey

University of Rochester Medical Center 601 Elmwood Avenue PO BOx 652 Rochester, NY 14642 USA Telephone: 585-366-5222 Email: jeffrey_rhodes@urmc.rochester.edu

Ricci, Michael A

Fletcher Allen Health Care 111 Colchester Ave., Patrick 226 Burlington, VT 05401 USA Telephone: 802-847-5155 Fax: 802-847-5907 Email: michael.ricci@vtmednet.org

Rich, Norman M (Lois)

USUHS/Department of Surgery 4301 Jones Bridge Road Bethesda, MD 20814 USA Telephone: 301-295-3155 Fax: 301-295-3627 Email: nrich@usuhs.mil

Ricotta, John J (Gloria)

SUNY at Stony Brook T19 HSC, Rm 020, Dept of Surgery Stony Brook, NY 11794-8191 USA Telephone: 631-444-7875 Fax: 631-444-8947 Email: jricotta@notes.cc.sunysb.edu

Rizvi, Adnan

Mayo Clinic Vascular Surgery 200 First St, SW Rochester, MN 55905 USA Telephone: 507-255-7062 Fax: 507-266-7156 Email: rizvi.adnan@mayo.edu

Robbins, Mark R (Leslie)

East Texas Vein Center 1028 E. Idel St., suite B Tyler, TX 75703 USA Telephone: 903-533-8720 Fax: 903-533-8720 Email: kids3md@msn.com

* Robicsek, Francis (Lilly)

Carolinas Heart Institute PO Box 32861 Charlotte, NC 28232-2861 USA Telephone: 704-355-4005 Fax: 704-355-6227 Email: frobicsek@sanger-clinic.com

Roddy, Sean P (Veronica)

The Vascular Group, PLLC 43 New Scotland Ave., MC157 Albany, NY 12208 USA Telephone: 518-262-8720 Fax: 518-262-6720 Email: roddys@albanyvascular.com

Rodman, Charles

Charles J. rodman MD, PA 740 Hospital Drive, Suite 150 Beaumont, TX 77701 USA Telephone: 409-832-8323 Fax: 409-832-4881 Email: crodmanmd@sbeglobal.net

Rodriguez, Agustin A (Liana Lopez)

University of Puerto Rico School of Medicine PO Box 364683 San Juan, PR 00936-4683 USA Telephone: 787-763-2440 Fax: 787-763-3898 Email: drgusrodriguez@aol.com

Rohrer, Michael J (Melody)

Univ. of TN Medical School 1325 Eastmoreland Ave., Ste. 310 Memphis, TN 38104 USA Telephone: 901-448-4100 Fax: 901-448-4110 Email: mrohrer@utmem.edu

* Rolley, Ronald T (Josette)

610 Ridgewood Dr West Lafayette, IN 47906 USA Fax: 317-477-9668

Rollins, David L (Carol)

3660 Euclid Ave, #107 Willoughby, OH 44094 USA Telephone: 440-269-8346 Fax: 440-975-5763 Email: dlrmd@safier.com

Rooke, Thom W (Julie)

Mayo Clinic 200 First St SW Rochester, MN 55905 USA Telephone: 507-266-7457 Fax: 507-266-1617 Email: rooke.thom@mayo.edu

Rosenfeld, Joel C (Beth)

St LUnited Kingdome's Hospital 801 Ostrum Street Bethlehem, PA 18015 USA Telephone: 610-954-2255 Fax: 610-954-6450 Email: rosenfj@slhn.org

Roupenian, Armen L

Vein & Laser Center NE Suite 305 45 Resnik Rd. Plymouth, MA 02360 USA Telephone: 508-747-1333 Fax: 508-747-2850 Email: whb8035@verizon.net

Rubin, Brian G

660 S. Euclid Ave., Campus Box 8109 St Louis, MO 63110-1094 USA Telephone: 314-362-7331 Fax: 314-362-7363 Email: rubinb@msnotes.wustl.edu

Rubin, Jeffrey R (Janis)

Detroit Medical Center/Harper Univ. Hospital Vascular Surgery 3990 John R Detroit, MI 48201 USA Telephone: 313-745-8637 Fax: 313-993-0244 Email: jrubin@med.wayne.edu

Ruby, Steven T (Gail)

St. Francis Hospital and Medical Center 1000 Asylum Ave, #2120 Hartford, CT 06105 USA Telephone: 860-246-4000 Fax: 860-527-6985 Email: vashartford@sbcglobal.net

* Rutherford, Robert B (Kay)

14337 Dorsal St Corpus Christie, TX 78418 USA Telephone: 361-949-0327 Fax: 361-949-8381 Email: rbruth@aol.com

Ryan, John J.

VÁ Medical Center 2501 East 22nd St. Sioux Falls, SD 57105 USA Telephone: 605-997-6277 Email: jjryan@usd.edu

* Sabety, Adrian M

The Cardiovascular Care Group 5 Franklin Ave Suite 310 Belleville, NJ 07109 USA Telephone: 973-759-9000

§ Sadick, Neil S

Sadick Aesthetic Surgery & Dermatology 911 Park Avenue New York, NY 10021-0337 USA Telephone: 212-772-7242 Fax: 212-517-9566 Email: nssderm@sadickdermatology.com

Sales, Clifford M (Kathy)

The Cardiovascular Care Group 5 Franklin Avenue, #310 Belleville, NJ 07109 USA Telephone: 973-759-9000 Fax: 973-751-3730 Email: csales@ thecardiovascularcaregroup.org

Salles-Cunha, Sergio X

Jobst Vascular Center 2109 Hughes Drive, 400 Toledo, OH 43606 USA Telephone: 419-291-2353 Fax: 419-479-6980

Salvian, Anthony J (Irene)

#1214-750 West Broadway Vancouver, BC V5Z 1J2 Canada Telephone: 604-874-0532 Fax: 604-874-7806 Email: salvian@pop.interchange.ubc.ca

Samson, Russell H

Mote Vascular Foundation 600 N. Cattlemen Road Suite 220 Sarasota, FL 34232-6422 USA Telephone: 941-371-6565 Fax: 941-377-7731 Email: rsamson@veinsandarteries.com

* Samuels, Peter B (Brenda)

2960 Neilson Way, Unit 502 Santa Monica, CA 90405-5373 USA Telephone: 310-396-5022 Fax: 310-450-8382

Schadeck, Michel P

Medical Center 5, rue Michel Chasles Paris F-75012 France Telephone: 33-1-43892220 Fax: 33-1-43896627 Email: flbskool@easynet.fr

Schanzer, Harry R (Helena)

Mount Sinai Médical Center 993 Park Avenue New York, NY 10028 USA Telephone: 212-396-1254 Fax: 212-396-1338 Email: harryschanzer@hotmail.com

Schellack, Jon V (Pamela)

Vascular Clinic 5425 Brittany Dr, Ste B Baton Rouge, LA 70808-4306 USA Telephone: 225-767-5479 Fax: 225-445-7202 Email: rsconyers@vasclin.com

§ Schepers, Helmut

Ganzoni Management AG St. Georgen Str. 70 Winterthur Zuerich 08401 Switzerland Telephone: 41-5-22650035 Fax: 41-5-22650001 Email: helmut.schepers@ganzoni.com

* Schmidt, Frank E (Donie)

1137 Jefferson Avenue New Orleans, LA 70115-3011 USA Telephone: 504-568-4576 Fax: 504-568-4633 Email: fesmd@bellsouth.net

Schneider, Joseph R (Shanda)

2650 Ridge Ave., Burch 100 Evanston, IL 60201 USA Telephone: 847-570-1009 Fax: 847-570-2899 Email: joe-schneider@northwestern.edu

* Schuler, James J (Catherine)

Univ of Illinois, Vasc Surg 1740 W Taylor, #2200 Chicago, IL 60612 USA Telephone: 312-996-7595 Fax: 312-996-2704 Email: mjmouw@uic.edu

Seabrook, Gary R (Nancy)

Medical College of Wisconsin 9200 W Wisconsin , Vasc Surgery MilwaUnited Kingdomee, WI 53226 USA Telephone: 414-456-8296 Fax: 414-456-6216 Email: gseabroo@mail.mcw.edu

* Segal Halperin, Boris M

Av Luis Maria Campos 1575, PB °C Buenos Aires 01426 Argentina Telephone: 541-784-9111 Fax: 541-784-9111 Email: borisegal@fibertel.com.ar

Shafique, Shoaib

Indiana University School of Medicine 1001 W. 10th Street OPE 303 Indianapolis, IN 46202 USA Telephone: 317-630-7879 Fax: 317-639-0271 Email: sshafiqu@iupui.edu

Shamma, Asad R (Lina)

Artery & Vein Institute P.O. Box 11-1666 Sodeco Sq; 8th Floor, Block B Beirut 111666 Lebanon Telephone: 961-375-0806 Email: shamuu@sovein.net

Saunders, Beverley

Charing Cross Hospital Fulham Palace Road Hammersmith London W6 United Kingdom Telephone: 20-8-8467335 Email: b.sharp@imperial.ac.United Kingdom

Shields, Raymond C (Opal)

Mayo Clinic 200 1st Street SW Rochester, MN 55905 USA Telephone: 507-266-9737 Fax: 507-266-1617 Email: shields.raymond@mayo.edu

Shortell, Cynthia K

DUnited Kingdome University Medical Center Box 3538 Durham, NC 27710 USA Telephone: 919-681-2915 Fax: 919-681-3563 Email: short018@mc.dUnited Kingdome.edu

Sidawy, Anton N (Mary)

7320 Yates Court McLean, VA 22101 USA Telephone: 202-745-8295 Fax: 202-745-8293 Email: ansidawy@aol.com

Silva, Michael B (Colleen)

TX Univ Health Sci Ctr, Dept Surg 3601 4th Street, Room 3A124 Lubbock, TX 79430 USA Telephone: 806-743-1306 Fax: 806-743-2359 Email: mbs2@aol.com

* Simonian, Simon J (Arpi)

3301 Woodburn Rd, #102 Annandale, VA 22003 USA Telephone: 703-573-5500 Fax: 703-573-3620 Email: veininstitute@netzero.net

§ Simons, Glen W

Kentucky Vein Care 125 East Maxwell, Suite 102 Lexington, KY 40508 USA Telephone: 859-455-8346 Fax: 859-455-8866 Email: gsimons@kyveincare.com

* Sladen, Joseph G (Jill)

3204 W. 26th Ave. Vancouver, BC V6L 1W1 Canada Telephone: 604-731-4085 Fax: 604-731-4081 Email: jsladen@interchange.ubc.ca

Sobel, Michael (Catherine)

VA Puget Sound Healthcare System 1660 S. Columbian Way, SS (112) Seattle, WA 98108-1597 USA Telephone: 206-764-2255 Fax: 206-764-2529 Email: michael.sobel@med.va.gov

Sottiurai, Vikrom S (Christine)

Center for Vein Health, Lutheran General Hospital 1775 Dempster St. Park Ridge, IL 60068 USA Telephone: 847-723-3008 Fax: 847-723-2535 Email: vikrom.sottiurai-md@ advocatehealth.com

Spence, Richard K (Claire)

St Agnes Medical Ctr 900 Caton Ave, #207, Dept Surgery Baltimore, MD 21229 USA Telephone: 410-368-2712 Fax: 410-951-4007 Email: rspence@stagnes.org

Stanley, Andrew C (Mary)

MCHV Campus Smith 111 Colchester Ave Burlington, VT 05401 USA Telephone: 802-656-8474 Fax: 802-656-0680 Email: andrew.stanley@uvm.edu

Steed, David L (Linda)

UPMC Shadyside 5200 Centre Avenue, Suite 307 Pittsburgh, PA 15232 USA Telephone: 412-623-8437 Fax: 412-623-8440 Email: steeddl@upmc.edu

Stonerock, Charles

1923 Brigadone Lane Florence, SC 29505-3241 USA Telephone: 843-676-2760 Fax: 843-067-62762 Email: therock.8@excite.com

Stoughton, Julianne (Mark N Nawrocki)

Vein Solutions 92 Montvale Ave, Ste #3200 Stoneham, MA 02180 USA Telephone: 781-438-8117 Fax: 781-438-8116 Email: doctor@veinsolutionsma.com

Suh, Bo Yang

Yeungnam Medical Center Dept. of Surgery 317-1 Daemyung-Dong, Nam-Gu Daegu 703-035 Korea Telephone: 82-5-36203583 Fax: 82-5-36241213 Email: bysuh@yumail.ac.kr

§ Sullivan, Cornelius A

Vasculart 200 Griffin Rd. Suite 6 Portsmouth, NH 03801 USA Telephone: 603-436-2002 Fax: 603-436-2006 Email: sullycamd@hotmail.com

* Sumner, David S (Martha)

2324 W. Lakeshore Drive Springfield, IL 62707 USA Telephone: 217-529-2910 Email: dsumner1@aol.com

* Taheri, Syde A (Rose Ann)

268 Dan Troy williamsville, NY 14221 USA Telephone: 716-633-1838 Fax: 716-634-4164 Email: staheri268@aol.com

Taylor, David C (Irene)

750 West Broadway, 708 Vancouver, BC V5Z 1H6 Canada Telephone: 604-876-2211 Fax: 604-874-7806 Email: dctaylor@interchange.ubc.ca

Thorpe, Patricia E

Venous Center 5 Woodland Heights Iowa City, IA 52240 USA Telephone: 319-688-5080 Fax: 319-688-5073 Email: patricia-thorpe@venous.com

Towne, Jonathan B (Sandra)

Medical College of Wisconsin 9200 West Wisconsin Ave MilwaUnited Kingdomee, WI 53226 USA Telephone: 414-456-6966 Fax: 414-456-6216 Email: jtowne@mail.mcw.edu

* Tretbar, Lawrence L

8787 Ballentine, 1200 Shawnee Mission, KS 66214 USA Telephone: 913-677-1776 Fax: 913-888-4081

Turnipseed, William D (Sandy)

Univ Wisconsin Clinical Sciences 600 North Highland Ave., G5-325 Madison, WI 53792 USA Telephone: 608-263-1388 Fax: 608-263-7652 Email: turnip@surgery.wisc.edu

Tzilinis, Argyrios

Anchor Vascular Surgery 800 Goodlette Rd. Suite 380 Naples, FL 34102 USA Telephone: 239-643-8794 Fax: 239-262-8129 Email: jtzilinis@hotmail.com

Valentin, Marlene D

Good Samaritan Hospital 375 Dixmyth Ave, Vascular Lab, Lvl 5 Cincinnati, OH 45220-2489 USA Telephone: 513-872-2769 Fax: 513-872-1549 Email: valentinmd@hotmail.com

Van Bemmelen, Paul S (Daphne)

Temple University 3401 No Broad St, Parkinson 4th Flr Philadelphia, PA 19140 USA Telephone: 215-707-3622 Fax: 215-707-5901 Email: vanbemp@tuhs.temple.edu

Varnagy, David

UMDNJ 150 Bergen Street Newark, NJ 07101 USA Telephone: 305-904-8149 Email: davidvarnagy@hotmail.com

Vasquez, Michael A (Melissa Ann)

SUNY Buffalo - Department of Surgery 415 Tremont Street Buffalo, NY 14120 USA Telephone: 716-690-2691 Fax: 716-690-2695 Email: mvasquezmd@roadrunner.com

Vazquez, Richard M

Northwestern Memorial Hospital 201 E. Huron St, Galter, Ste 11-250 Chicago, IL 60611 USA Telephone: 312-649-6562 Fax: 312-649-9027 Email: dry@veincare.com

Vedantham, Suresh

Mallinckrodt Institute of Radiology 510 S. Kings Highway Blvd. Box 8131 St. Louis, MO 63110 USA Telephone: 314-719-3431 Fax: 314-362-2276 Email: vedanthams@mir.wustl.edu

Villavicencio, J. Leonel (Susy) USUHS, Prof Surgery 4301 Jones Bridge Rd Bethesda, MD 20814 USA Telephone: 202-782-6592 Fax: 202-782-3371

Email: jvillavicencio@usuhs.mil Wakefield, Thomas W (Mary)

Univ of Michigan Medical Ctr 1500 E Medical Ctr Dr, THCC 2210 Ann Arbor, MI 48109-0329 USA Telephone: 734-936-5820 Fax: 734-647-9867 Email: thomasww@umich.edu

Walsh, Daniel B (Teri)

Dartmouth-Hitchcock Med Ctr One Medical Center Dr Lebanon, NH 03756 USA Telephone: 603-650-8191 Fax: 603-650-4973 Email: daniel.walsh@hitchcock.org

Wasserman, Dean H (Regina)

Vein Treatment Ctr of NJ 1 West Ridgewood Ave Paramus, NJ 07652 USA Telephone: 201-612-1750 Fax: 201-612-1760 Email: cutter2d@aol.com

Webster, Marshall W (Bonnie)

Univ of Pittsburgh Medical Center 200 Lothrop St, #9019 Forbes Tower Pittsburgh, PA 15213 USA Telephone: 412-647-1912 Fax: 412-647-1919 Email: webstermw@msx.upmc.edu

Weingarten, Michael S (Carol)

Drexel University College of Medicine / Hahnemann Hospital 245 N. 15th Street# 7150 Mailstop 413 Philadelphia, PA 19102 USA Telephone: 215-762-4005 Fax: 215-762-8699 Email: michael.weingarten@drexelmed.edu

Weiss, Robert A (Margaret)

Aspen Mill Professional Bldg 54 Scott Adam Rd, #301 Hunt Valley, MD 21030 USA Telephone: 410-666-3960 Fax: 410-666-3981 Email: ksorenson@mdlaserskinvein.com

Welch, Harold J (Cynthia)

Lahey Clinic 41 Mall Rd, Peripheral Vasc Surgery Burlington, MA 01805 USA Telephone: 781-744-8193 Fax: 781-744-5744 Email: harold.j.welch@lahey.org

Wennberg, Paul W (Julie)

Mayo Clinic 200 First Street SW Rochester, MN 55905 USA Telephone: 507-266-7231 Fax: 507-266-1617 Email: wennberg.paul@mayo.edu

* Wheeler, H. Brownell (Betty)

Univ of Mass Medical School 55 Lake Ave North, #S3-810, Surgery Worcester, MA 01655 USA Telephone: 508-856-2201 Fax: 508-856-6941 Email:

* Williams, G. Melville (Linda) Johns Hopkins Hospital 600 No Wolfe St, Harvey 611 Baltimore, MD 21287-8611 USA Telephone: 410-955-5165 Fax: 410-614-2079 Email: melwill@erols.com

Williams, David

University of Michigan B1-D530 1500 E. Medical Center Drive Ann Arbor, MI 48109-0030 USA Telephone: 734-662-2717 Email: davidwms@med.umich.edu

Wolk, Seth W (Ruthanne)

Restoration Vein Care 5333 McAuley Dr., Suite 4016 Ann Arbor, MI 48106 USA Telephone: 734-712-4310 Email: wolksw@trinity-health.org

Yamaki, Takashi

Tokyo Women's Medical University 8-1, Kawada-cho, ShinjUnited Kingdomu-ku Tokyo 162-8666 Japan Email: yamaki@prs.twmu.ac.jp

* Yao, James S. T. (Louise)

Northwestern University Med. School 201 East Huron St, #10-105 Chicago, IL 60611 USA Telephone: 312-695-2716 Fax: 312-695-4955 Email: jyao@nmh.org

Yellin, Albert E (Elissa)
 59-415 Kawowo Road
 Haleiwa, HI 96712
 USA
 Email: aeyellin@hawaii.rr.com

Yunus, Tahir

William Beaumont Hospital 3601 W. 13 Mile Road Royal Oak, MI 48073 USA Telephone: 248-854-7972 Email: tahirey@yahoo.com

§ Zakaria, Aamir M

SIU School of Medicine PO box 19638 Springfield, IL 62794-9638 USA Telephone: 217-545-8444 Fax: 217-545-2563 Email: azakaria@siumed.edu

Zatina, Michael A (Katie)

3350 Wilkens Ave, Ste #100 Baltimore, MD 21229-4615 USA Telephone: 410-368-2712 Fax: 410-368-3569 Email: mzatina@stagnes.org

Zayyat, Elie J (Tracy)

Good Samaritan Hospital 375 Dixmythave-Med Edu 3rd Fl Cincinnati, OH 45220 USA Telephone: 513-844-1000 Fax: 513-895-1254 Email: etzayyat@aol.com

Zierler, Brenda K

University of Washington 1959 NE Pacific St, Box 357266 Seattle, WA 98195-7266 USA Telephone: 206-616-1910 Fax: 206-616-7495 Email: brendaz@u.washington.edu

Zierler, R. Eugene

University of Washington 1959 NE Pacific Street Box 356410 Seattle, WA 98195 USA Telephone: 206-543-3095 Fax: 206-616-7495 Email: gzierler@u.washington.edu

Zimmet, Steven

Chairman, ACP Foundation 1500 West 34th Street Austin, TX 78703 USA Telephone: 512-485-7700 Email: zimmet@skin-vein.com

Zubicoa, Santiago Ezpeleta

Hospital Ruber Internacional c/ la Maso N. 38 Madrid 28034 Spain Telephone: 34-9-13875157 Fax: 34-9-13875158 Email: ana.b.romero@aexp.com

INTERNATIONAL MEMBERS

Arfvidsson, Berndt

University Hospital of Orebro Orebro 70185 Sweden Telephone: 46-1-9125439 Fax: 46-1-9125439 Email: berndt.arfvidsson@orebroll.se

Balas, Panayiotis E

Hiraclitou 4 Athens GR-1067 Greece Telephone: 30-1-6712055 Fax: 30-1-6712055 Email: balasgr@compulink.gr

Bass, Arie

Assaf Harofeh Medical Ctr. Dept. of Vascular Surgery Zerifin 70300 Israel Email: arbas@post.tau.ac.il

Carpentier, Patrick H

Grenoble University Hospital Vascular Medicine Clinic Grenoble F38043 France Telephone: 33-4-76768735 Fax: 33-4-76768735 Email: patrick.h.carpentier@orange.fr

Christenson, Jan T (Suzy)

University of Geneva, Dept. Cardiovascu 24 rue Micheli-du-Crest Geneva CH-1292 Switzerland Telephone: 41-2-23727634 Fax: 41-2-23727634 Email: jan.christenson@hcuge.ch

Cigorraga, Jorge Raul (Maria Isabel Trapaglia)

Av Las Heras 2223 5" A Buenos Aires 1425 Argentina

Cornu-Thenard, Andre M

Saint Antoine Hospital 113 avenue Charles de Gaulle Neuilly / Seine 92200 France Telephone: 33-1-47451421 Fax: 33-1-47451421 Email: andre.cornuthenard@ wanadoo.fr

Davies, Alun Huw

Charing Cross Hospital Fulham Palace Rd, Surgery, 4th Floor London W6 8RF United Kingdom Telephone: 44-2-088467362 Fax: 44-2-088467362 Email: a.h.davies@ic.ac.uk

di Marzo, Luca

Department of Surgery "Pietro Valdoni" University of Rome "La Sapienza" Viale del Policlinico, 155 Rome 161 Italy Telephone: 39-0-649972203 Fax: 39-0-649972203 Email: luca.dimarzo@uniroma1.it

Disselhoff, Ben

Mesos Medical Center Dept. of Vascular Surgery 8605 RP Utrecht Utrecht 3527CE Netherlands Email: bcvmdisselhoff@mesos.nl

Farmache, Alejandro H (Rosa Imes)

Instituto de Flebolog a Necochea 350 1 Piso Dpto 12 Ciudad Mendoza 5500 Argentina Telephone: 54-2-614210997 Fax: 54-2-614210997 Email: afarmache@speedy.com.ar

Fegan, William G

PO Box 100 Lamu Kenya

Guex, Jean-Jerome (Genevieve)

Angiology Clinic 32, Boulevard Dubouchage Nice F-06000 France Telephone: 33-4-93854130 Fax: 33-4-93854130 Email: jj.guex@wanadoo.fr

Gupta, Prem C (Laxmi)

Medwin Hospital #311 Maruti Sadan, 6-3-1117 Begumpet Hyderbad 500-016 India Telephone: 91-4-023201120 Fax: 91-4-023201120 Email: pcgupta10@hotmail.com

Hartung, Olivier

Service de Chirurgie Vasculaire, CHU Nord Chemin des Bourrelys Marseille 13015 France Telephone: 33-4-91968370 Fax: 33-4-91968370 Email: olivier.hartung@ap-hm.fr

Hoshino, Shunichi (Hiroko)

Fukushima Daiichi Hospital 16-2 Nariide, Kitasawamata Fukushima-city 960-8251 Japan Telephone: 11-8-1245575064 Fax: 11-8-1245575064 Email: shurhoshino@r9.dion.ne.jp

Ishimaru, Shin

Tokyo Med College, Surgery 6-7-1 Nishi-shinjuku, Shinjuku-ku Tokyo 160-0023 Japan Telephone: 81-3-33422827 Fax: 81-3-33422827 Email: ishimaru@tokyo-med.ac.jp

Kim, Young-Wook (SeonMin Park)

Samsung Medical Center 50, Ilwon-Dong, Gangnam-Gu Seoul 135-710 Korea Telephone: 82-2-34100040 Fax: 82-2-34100040 Email: ywkim@smc.samsung.co.kr

Komlos, Pedro P

Pedro Pablo Komlos Vas Surg Clinic rua Dr Florencio Ygartua St, 131rm605 Porto Alegre- RS 90430-010 Brazil Telephone: 55-5-133302315 Fax: 55-5-133302315 Email: ppkomlos@terra.com.br

Kompf, Boguslaw

Klinika Zdrowych Nog ul. Reduty Ordona 54/1 71-202 Szczecin Poland Email: dr@kompf.com

Liew, Ngoh C

University of Putra Malaysia Dept. of Surgery Kuala Lumpor 50586 Malaysia Telephone: 60-3-20501076 Fax: 60-3-20501076 Email: liewnc@yahoo.com

Matsubara, Junichi (Junko)

Kanazawa Med Univ, 1-1 Daigaku Uchinada-Machi, Kahoku-gun Ishikawa-Ken 920-02 Japan Telephone: 81-0-762862322 Fax: 81-0-762862322 Email: matsujun@kanazawa-med.ac.jp

Milleret, Rene

Vein Center 2 Rue de Verdun Montpellier 34000 France Email: rmilleret001@rss.fr

Ogawa, Tomohiro

CV Disease Ctr/Fukushima Daiichi Hosp 16-2 Kitasawamata Nariide Fukushima City 960-8251 Japan Telephone: 81-2-45575064 Fax: 81-2-45575064 Email: tomo-ogawa@msb.biglobe.ne.jp

Osse, Francisco

Venaclinic Rua Lomas Valentinas, 278 Sao Paulo 05084-010 Brazil Telephone: 55-1-138359365 Fax: 55-1-138359365 Email: fjosse@uol.com.br

Papendieck, C. M. (Laura)

Universidad del Salvador Catamarca 3179 - 1636 Olivos Buenos Aires 1636 Argentina Telephone: 54-1-147990740 Fax: 54-1-147990740 Email: cpapen@intramed.net.ar

Pietravallo, Antonio F. R.

Inst Privado de Flebologia Av Callao 1243, 10 B Buenos Aires 1023 Argentina Telephone: 54-1-8144496 Fax: 54-1-8144496 Email: flebologiapietravallo@hotmail.com

Rasmussen, Lars H (Birgit)

Kirurgisk Center Naestved Eskadronvej 4 A Naestved 4700 Denmark Telephone: 45-5-5723038 Fax: 45-5-5723038 Email: Ihr@varix.dk

Richardson, Graeme D (Dianne)

Rural Clinical School,UNSW PO Box 5695 Wagga Wagga 2650 Australia Email: richo2@aapt.net.au

Sakuda, Hitoshi

Second Department of Surgery, University of the Ryukyus 207 Uehara, 2nd Dept of Surgery Nishihara Okinawa 903-0215 Japan Telephone: 81-9-88951422 Fax: 81-9-88951422 Email: sakuda-h@med.u-ryukyu.ac.jp

Schapira, Armando E (Estela)

Clinica de Flebolinfologia Buenos Aires 1013 Rosario 2000 Argentina Telephone: 54-3-414242634 Fax: 54-3-414242634 Email: schapira@cimero.org.ar

Schultz-Ehrenburg, Ulrich (Helga neeDehnke)

Berlin-Weissensee Health Ctr Schonstr 5, Dermatology & Phlebology Berlin 13086 Germany Telephone: 49-3-305622654 Fax: 49-3-305622654 Email: u.schultz-ehrenburg@t-online.de

Scurr, John H

Lister Hospital, Lister House Chelsea Bridge Rd London SW1W 8RH United Kingdom Telephone: 44-0-2702599938 Fax: 44-0-2702599938 Email: jscurr@uk-consultants.co.uk

Shaidakov, Evgeny V

Military Medical Acadamy Fontanka 106 St. Petersburg 198013 Russia Email: shaidak@mail.wplus.net

Simkin, Roberto

University of Buenos Aires Argentina Talcahuano 1155, P. Baja Dto.5 Buenos Aires 1013 Argentina Telephone: 54-1-148054774 Fax: 54-1-148054774 Email: robsim@ciudad.com.ar

Uhl, Jean-Francois

Vanuse Veins Surgical Center 113 Av ch de Gaulle Neuilly-sur-seine 92200 France Telephone: 33-1-47472060 Fax: 33-1-47472060 Email: jf.uhl@wanadoo.fr

Vandendriessche-Hobbs, Marianne

Vein Clinic 288 Maaltebrugge St Gent B9000 Belgium Telephone: 32-9-2200781 Fax: 32-9-2200781 Email: mvandendriessche@hotmail.com

Wittens, Cees H. A. (Janny)

hagaziekenhuis, Den Haag, Netherlands Bergse Linker Rottekade 204 Rotterdam 3056 LE Netherlands Telephone: 31-1-04616769 Fax: 31-1-04616769 Email: ceeswittens@chello.nl

Zamboni, Paolo (Elena)

Univ Degli Studi Di Ferrara 203 Corso Giovecca, Surgery Ferrara 44100 Italy Telephone: 39-0-532237443 Fax: 39-0-532237443 Email: cfr@unife.it AMERICAN VENOUS FORUM

Geographical Roster

ALABAMA

Birmingham Passman, Marc A

ARKANSAS

Little Rock Ferris, Ernest J

ARIZONA

Scottsdale Morrison, Nick

CALIFORNIA

Agoura Barker, Wiley F

Baldwin Park Murray, James D

Beverly Hills Gradman, Wayne S

Dana Point Cannon, Jack A

Escondido Bulkin, Anatoly

Fresno Elmore, Frederick A

Irvine Kanter, Alan

La Jolla Bergan, John J Cheng, Van Le Delaria, Giacomo A Fronek, Arnost OByrne, Margaret G Pascarella, Luigi

Loma Linda Hasaniya, Nahidh W

Newport Beach Arata, Michael Orange Flanigan, D. Preston

Panorama City Cerveira, Joaquim J

Portola Valley Fogarty, Thomas J

Rancho Palos Verdes Donayre, Carlos E

Roseville Monahan, Daniel L.

San Diego Angle, Niren

San Francisco Denbo, Howard E Long, John B

San Mateo Harris, Edmund J

Santa Barbara Mckittrick, James E

Santa Monica Samuels, Peter B

Seal Beach Gaspar, Max R

Stanford Harris, E. John

Torrance Duffy, David M

Walnut Creek Isaacs, Mark

COLORADO

Aurora Hammond, Sharon L

Fort Collins Kaufman, Steven L

Palisade Bernhard, Victor M

CONNECTICUT

Farmington Menzoian, James O

Greenwich Mulcare, Robert

Hartford Ruby, Steven T

DISTRICT OF COLUMBIA

Washington Beavers, Frederick P Depalma, Ralph G Feied, Craig F Giordano, Joseph M Laredo, James

DELEWARE

Newark Gomes, Mario N

Wilmington Kerstein, Morris D

FLORIDA

Boca Raton Meretei, Attila

Jacksonville Hakaim, Albert G

Miami

Almeida, Jose Ignacio Ginzburg, Enrique Kang, Steven S

Naples Tzilinis, Argyrios

Palm Beach Gardens Miskin, Barry M

Sarasota Samson, Russell H

Tampa Kerr, Thomas M

West Palm Beach Lynn, Richard A

Weston Fernandez, Bernardo B

GEORGIA

Atlanta Ferrier, Frank Kasirajan, Karthikeshwar

Gainesville Procter, Charles D

Savannah Alpert, Joseph

HAWAII

Haleiwa Yellin, Albert E

Honolulu Kistner, Robert L Lurie, Fedor Masuda, Elna M

IOWA

lowa City Thorpe, MD, Patricia E

West Des Moines Anderson, Robert

ILLINOIS

Arlington Heights Forrestal, Mark

Chicago Bassiouny, Hisham S Durham, Joseph R Matsumura, Jon S McCarthy, Walter J Morasch, Mark D Pearce, William H Schuler, James J Vazquez, Richard M Yao, James S. T

Evanston Schneider, Joseph R

LaGrange Gocke, John

Maywood Leon, Luis

Park Ridge Buckman, Jeffrey Sottiurai, Vikrom S Skokie Caprini, Joseph A

Springfield McLafferty, Robert B Sumner, David S Zakaria, Aamir M

INDIANNA

Carmel Finkelmeier, William R

Indianapolis Dalsing, Michael C Goodson, Spencer F Lalka, Stephen G Shafique, Shoaib

West Lafayette Rolley, Ronald T

KANSAS

Shawnee Tretbar, Lawrence L

KENTUCKY

Lexington Simons, Glen W

Pikeville Collins, David E.

LOUISIANNA

Baton Rouge Frusha, John D Schellack, Jon V

New Orleans Hollier, Larry H Schmidt, Frank E

Shreveport Phifer, Travis J

MASSACHUSETTS

Arlington^e Flynn, William F Boston Abbott, William M Baldwin, John C Cantelmo, Nancy L lafrati, Mark D O'Donnell, Thomas F Polak, Joseph F

Brighton Razvi, Syed A

Burlington Welch, Harold J

Framingham Donaldson, Magruder C

Plymouth Roupenian, Armen L

Stoneham Stoughton, Julianne

Wellesley Persson, Alfred V

West Roxbury Raffetto, Joseph D

Westport Carney, Wilfred I

Worcester Messina, Louis M Wheeler, H. Brownell

MARYLAND

Baltimore Flinn, William R Spence, Richard K Williams, G. Melville Zatina, Michael A

Bethesda Rich, Norman M Villavicencio, J. Leonel

Hunt Valley Weiss, Robert A

Potomac Gillespie, David L

Towson Buchbinder, Dale

MAINE

Portland Eldrup-Jorgensen, Jens

MICHIGAN

Ann Arbor Greenfield, Lazar J Henke, Peter K Myers, Jr., Daniel Wakefield, Thomas W Williams, David Wolk, Seth W

Bingham Farms Brown, O. William

Detroit

Mattos, Mark A Nypaver, Timothy J Rubin, Jeffrey R

Grand Rapids Mansour, M. Ashraf

Livonia Pfeifer, John R

Petoskey Kazmers, Andris

Royal Oak Yunus, Tahir

Troy Engle, Jennifer S

West Bloomfield Elliott, Joseph P Granke, Kenneth

MINNESOTA

Rochester

Agarwal, Gautam Bjarnason, Haraldur Felty, Cindy Gloviczki, Peter Kalra, Manju Knott, Andrew W Lall, Purandath Lofgren, Eric P Lofgren, Karl A Noel, Audra A Oderich, Gustavo Rizvi, Adnan Rooke, Thom W Shields, Raymond C Wennberg, Paul W

MISSISSIPPI

Flowood Neglen, Peter Raju, Seshadri

MISSOURI

Columbia Gardner, Glenn P

St Louis Binnington, H. Bradley Kloecker, Richard J Rubin, Brian G Vedantham, Suresh

NORTH CAROLINA

Chapel Hill Marston, William A

Charlotte Robicsek, Francis

Durham O'Shea, Susan I Shortell, Cynthia K

Greenville Powell, C. Steven

NEBRASKA

Omaha Lynch, Thomas G

NEW HAMPSHIRE

Lebanon Beebe, Hugh G Walsh, Daniel B

Manchester Furey, Patricia C

Portsmouth Sullivan, Cornelius A

NEW JERSEY

Belleville Sabety, Adrian M Sales, Clifford M

Denville Araki, Clifford T Mintz, Bruce Englewood Elias, Steven

Highland Park Konigsberg, Stephen F

Morristown Moritz, Mark W Oliver, Mark A

New Brunswick Haser, Paul B

Newark

Abai, Babak Hobson, Robert W Jamil, Zafar Labropoulos, Nicos Lal, Brajesh K Padberg, Frank T Pappas, Peter J Varnagy, David

Paramus Wasserman, Dean H

Somerset Deak, Steven T

NEW MEXICO

Albuquerque Corson, John D

Clovis Paladugu, Ramesh

Santa Fe Martin, Alfred J

NEVADA

Reno Daake, John W Merchant, Robert F

NEW YORK

Albany

Chang, Benjamin B Darling, R. Clement Roddy, Sean P

Brooklyn

Ascher, Enrico Hingorani, Anil P Mutyala, Manikyam Puggioni, Alessandra Rai, Dinker B Buffalo Harris, Linda M

Vasquez, Michael A

New York

Adelman, Mark A Baron, Howard C Gagne, Paul Green, Richard M Kabnick, Lowell S Kent, K. Craig Min, Robert J Olin, Jeffrey W Sadick, Neil S Schanzer, Harry R

Rochester Deweese, James A

Illig, Karl A Rhodes, Jeffrey

Roslyn Chang, John B

Schenectady Blumenberg, Robert M

Staten Island Fodera, Maria Elena

Stony Brook Criado, Enrique Gasparis, Antonios P Ricotta, John J

Williamsville Taheri, Syde A

OHIO

Cincinnati Cranley, Robert D Hutto, John D. Kempczinski, Richard Lohr, Joann M

Muck, Patrick E Pringle, Timothy C Reed, Amy B Valentin, Marlene D Zayyat, Elie J

Cleveland Carman, Teresa L

Columbus Franz, Randall

Dayton Lemmon, Gary W Steubenville Ramnauth, Subhash C

Toledo

Balkany, Louis Comerota, Anthony J Dosick, Steven M Gale, Steven S

Kritpracha, Boonprasit Nazzal, Munier M.S. Salles-Cunha, Sergio X

Westlake Navarro, Felipe

Willoughby Rollins, David L

OREGON

Portland Edwards, James M Moneta, Gregory L Pavcnik, Dusan

PENNSYLVANIA

Bethlehem Rosenfeld, Joel C

Easton Balshi, James D Fisher, Jay B

Mechanicsburg Calcagno, David

Philadelphia

Blebea, John Calligaro, Keith D Merli, Geno J Van Bemmelen, Paul S Weingarten, Michael S Cho, Jae-Sung

Pittsburgh Jarrett, Fredric Steed, David L Webster, Marshall W

Wayne Ernst, Calvin B

Wilkesbarre Gruneiro, Laura A

York Castronuovo, John J

PUERTO RICO

San Juan Rodriguez, Agustin A

RHODE ISLAND

Providence Patterson, Robert B

SOUTH CAROLINA

Aiken Ratliff, Jack L

Charleston Hallett, John W

Florence Stonerock, Charles

SOUTH DAKOTA

Sioux Falls Ryan, John J.

TENNESSEE

Knoxville Goldman, Mitchell H

Memphis Rohrer, Michael J

TEXAS

Austin Dilling, Emery Zimmet, Steven

Beaumont Rodman, Charles

Corpus Christie Rutherford, Robert B

Dallas Clagett, G. Patrick

Galveston Hunter, Glenn C Killewich, Lois A Pounds, Lori C

Houston Bush, Ruth Lin, Peter Lumsden, Alan B Peden, Eric Lubbock Silva, Michael B

McKinney Proctor, Mary C

San Antonio Martinez, Jeffrey M.

Temple Bohannon, W. Todd

Tyler Robbins, Mark R

Victoria Johnston, Robert H

VIRGINIA

Alexandria Cordts, Paul R

Annandale Simonian, Simon J

Charlottesville Cherry, Kenneth J Owens, Lewis

McLean Sidawy, Anton N

Norfolk Bonawitz, Cara A

Portsmouth Arbid, Elias J

Reston Lee, Byung-Boong

Williamsburg Delaurentis, Dominic A

VERMONT

Burlington Ricci, Michael A Stanley, Andrew C

WASHINGTON

Seattle

Meissner, Mark H Sobel, Michael Zierler, Brenda K Zierler, R. Eugene

Vancouver

Nicholls, Stephen

WISCONSIN

Madison Carr, Sandra C Turnipseed, William D

Manitowoc Gueldner, Terry L.

Milwaukee

Brown, Kellie Cambria, Robert A Mewissen, Mark W Seabrook, Gary R Towne, Jonathan B

WEST VIRGINIA

Charleston AbuRahma, Ali F Boland, James P

INTERNATIONAL MEMBERS

ARGENTINA

Buenos Aires

Cigorraga, Jorge Raul Enrici, Ermenegildo A Papendieck, C. M. Pietravallo, Antonio F. R. Segal Halperin, Boris M Simkin, Roberto

Mendoza Farmache, Alejandro H

Rosario Schapira, Armando E

AUSTRALIA

Wagga Wagga Richardson, Graeme D

AUSTRIA

Vienna Partsch, Hugo

BELGIUM

Gent Vandendriessche-Hobbs, Marianne

BRAZIL

Porto Alegre-RS Komlos, Pedro P

Sao Paulo Osse, Francisco

CANADA

Calgary Hill, Douglas

Hamilton Hirsh, Jack

Quebec Dion, Yves M

Vancouver Salvian, Anthony J

Sladen, Joseph G Taylor, David C CYPRUS

Nicosia Nicolaides, Andrew N

DENMARK

Naestved Rasmussen, Lars H

FRANCE

Cagnes sur ner Pittaluga, Paul

Chassieu Perrin, Michel

Grenoble Carpentier, Patrick H

Marseille Hartung, Olivier

Montpellier Milleret, Rene

Neuilly/Seine Cornu-Thenard, Andre M

Neuilly-sur-seine Uhl, Jean-Francois

Nice Guex, Jean-Jerome

Paris Cazaubon, Nichele Natali, Jean P Schadeck, Michel P

GERMANY

Berlin Schultz-Ehrenburg, Ulrich

Bonn Rabe, Eberhard

Heidelberg Proebstle, Thomas

Kassel Gruss, Jorg D

Nuremberg Noppeney, Thomas

GREECE

Athens Balas, Panayiotis E Liasis, Nikolaos E.

INDIA

Hyderbad Gupta, Prem C

ISRAEL

Zerifin Bass, Arie

ITALY

Ferrara Zamboni, Paolo

Rome Allegra, Claudio Caggiati, Alberto di Marzo, Luca

JAPAN

Fukushima City Ogawa, Tomohiro Hoshino, Shunichi

Ishikawa-Ken Matsubara, Junichi

Nishihara Okinawa Sakuda, Hitoshi

Tokyo Ishimaru, Shin Yamaki, Takashi

KENYA

Lamu Fegan, William G

KOREA

Daegu Suh, Bo Yang

KOREA

Seoul Kim, Young-Wook

LEBANON

Beirut Shamma, Asad R

LUXEMBOURG

Goetzingen Lamesch, Alfred J

MALAYSIA

Kuala Lumpor Liew, Ngoh C

MEXICO

Mexico City Paramo-Diaz, Marcelo

NETHERLANDS

Rotterdam Wittens, Cees H. A

Utrecht Disselhoff, Ben

POLAND

Szczecin Kompf, Boguslaw

RUSSIA

St. Petersburg Shaidakov, Evgeny V

SPAIN

Madrid Monedero, Javier Leal Zubicoa, Santiago Ezpeleta

SWEDEN

Helsingborg Eklof, Bo G

Linkoping Thulesius, Olav

Orebro Arfvidsson, Berndt

Uppsala Bergqvist, David

SWITZERLAND

Geneva

Christenson, Jan T

Lucerne

Lauber, Andre F

Strafa Bollinger, Alfred

Zuerich

Schepers, Helmut

TURKEY

Istanbul Kurtoglu, Mehmet H.

UNITED KINGDOM

Channel Islands

Browse, Norman L

Edinburgh

Ruckley, C. Vaughan

London

Burnand, Kevin G Davies, Alun Huw Hobbs, John T Saunders, Beverley Scurr, John H

Solihull

Bradbury, Andrew W

Wexham

Coleridge Smith, Philip D

WEST INDIES

Trinidad

Maharaj, Dale A

THE AMERICAN VENOUS FORUM

BY-LAWS

ARTICLE I - NAME

The name of this organization shall be **THE AMERICAN VENOUS FORUM**.

ARTICLE II - OBJECTIVES

The objectives of this organization shall be (1) to promote the study of or research in venous diseases; (2) to contribute to the active continuing education of its membership; (3) to hold annual meetings; and (4) to encourage the development and dissemination of knowledge regarding venous disease.

Notwithstanding the foregoing, (a) no part of the organization's net earnings or assets shall inure to the benefit of any member, officer, or other person, except that the organization shall be authorized and empowered to pay reasonable compensation for services rendered and to make other payments and distributions in furtherance of the purposes set forth above, and (b) the organization shall not carry on any activity not permitted to an organization exempt from Federal income tax under Section 501(c)(6) of the Internal Revenue Code of 1954, as amended (the "Code") or the corresponding provision of any future United States revenue statute.

ARTICLE III - MEMBERSHIP

Membership in the Venous Forum may include any physicians certified by their respective specialty Certifying Boards in the applicant's Country of practice who have demonstrated an interest in and contribution to the management of venous problems and who are in good standing in their State or Provincial Medical Societies. From time to time, the Membership Committee may recommend membership to scientists who are not M.D.'s and/or do not possess a doctoral degree but have demonstrated a major commitment to issues of venous disease.

- 1. Active Members: as identified above. Active members shall pay dues and have full voting privileges. Attendance at the Annual Scientific Program shall be expected of all Active members.
- 2. Senior Members: included will be active members who have reached the age of 65 years; or members for whom, for reasons of health or other just cause, the Executive Committee recommends this category. They shall not be bound by meeting attendance and dues may be waived upon written request by Senior Member to waive dues. The Executive Committee may approve or disapprove the request at an executive meeting.
- 3. Honorary Members: individuals who have made outstanding contributions in the field of venous science. They shall not pay dues nor shall they have voting privileges.
- 4. Associate Members: Individuals who have an interest in the management of venous disorders, but do not necessarily hold a doctoral degree, such as nurses, registered vascular technologists, etc. Associate members will pay membership dues determined by the Executive Committee. Associate members are not eligible to vote or hold elective office.

5. Candidate Members: Physicians who are currently serving in a capacity of a resident or fellow in a post-doctoral training program and have demonstrated interest in and have made a contribution to the management of venous disease. Candidate members are not eligible to vote or hold elective office and are required to pay membership dues as set by the Executive Committee. Membership in this category shall not exceed 3 years. At the conclusion of post-doctoral training, Candidates may opt to become Active Members, by notifying the Forum in writing. In this instance, the application process will be waived, and the name shall automatically be placed on the Ballot.

ARTICLE IV - ELECTION OF MEMBERS

- 1. The process of election of ACTIVE members of the Society shall be as follows:
 - a. Applications must be accompanied by letters from the sponsor and two endorsers all of whom should be members of the American Venous Forum.
 - b. Application forms presenting the curricula vitae of the candidates and signed by them shall be in the hands of the Secretary before the executive session at which it is desired that the candidate be considered for election.
 - c. The Secretary shall send to the Chairman of the Membership Committee these applications with all pertinent data before the annual meeting. The Membership Committee shall review the professional qualifications of the candidates.
 - d. The Chairman of the Membership Committee shall meet with the Executive Committee for the purpose of presenting the recommendations of the Membership Committee.
 - e. The names of the candidates recommended by the Executive Committee for election shall be submitted by the Secretary to the membership in his or her annual report.
 - f. Election to membership shall be by secret ballot, by a three fourths affirmative vote of those members present and voting at the Annual Business Meeting.
 - g. A candidate who fails of election at one meeting may be presented to the membership at the next two (2) annual meeting of the Forum. The name of a candidate who fails of election a third time shall be dropped from the list of applications for membership. Such candidate's application may be resubmitted after an interval of two (2) years.
 - h. New Member Attendance: Candidates, following their election to membership at the Annual Business Meeting of the organization, will be required to attend the next Annual Meeting of the Forum to be formally introduced to the membership.
- 2. The process of election for Associate and Candidate Members shall be as follows:
 - a. Application forms presenting the curricula vitae of the candidates and signed by them shall be in the hands of the Secretary before the executive session at which it is desired that the candidate be considered for election.
 - b. The Secretary shall send to the Chairman of the Membership Committee these applications with all pertinent data before the annual meeting. The Membership Committee shall review the professional qualifications of the candidates.
 - c. The Chairman of the Membership Committee shall meet with the Executive Committee for the purpose of presenting the recommendations of the Membership Committee.

- d. The names of the candidates recommended by the Executive Committee for election shall be submitted by the Secretary to the membership in his or her annual report.
- e. Election to membership shall be by secret ballot, by a three fourths affirmative vote of those members present and voting at the annual business meeting
- f. A candidate who fails of election at one meeting may be presented to the membership at the next two (2) annual meeting of the Forum. The name of a candidate who fails of election a third time shall be dropped from the list of applications for membership. Such candidate's application may be resubmitted after an interval of two (2) years.
- g. New Member Attendance: Candidates, following their election to membership at the Annual Business Meeting of the organization, will be required to attend the next Annual Meeting of the Forum to be formally introduced to the membership.
- 3. The process of election for Honorary Members of the Forum shall be as follows:
 - a. Any Active or Senior member may nominate an individual for Honorary membership. The name and a brief description of the accomplishments of the nominee must be submitted to the Secretary before the Executive Session at which it is desired the nominee be considered for honorary membership. The Secretary shall distribute this information to the Honorary Membership Committee consisting of three (3) immediate past Presidents of the Executive Committee before the annual meeting.
 - b. The Honorary Membership Committee shall make its recommendations to the Executive Committee.
 - c. Following its deliberation, the Executive Committee may recommend that the candidate's name be submitted by the Secretary to the membership in the annual report at the Annual Business Meeting of the Forum.
 - d. Election to Honorary Membership shall be by secret ballot by three fourths affirmative vote of the membership present and voting at the Annual Business Meeting.

ARTICLE V - EXECUTIVE COMMITTEE

- 1. The Executive Committee of the Forum shall direct its activities.
- 2. The Executive Committee shall be composed of the President, the President Elect, the Secretary, the Treasurer, the Recorder, three Councilors and the immediate three Past Presidents and the Archivist.
- 3. The Executive Committee shall be the governing body of the Forum and shall have full power to manage and act on all affairs on the Forum except as follows:
 - a. It may not, without the approval of the Forum membership at an annual executive session, alter the initiation fees or levy any assessment against the membership, except that it may, set the annual dues rates and, in individual cases, waive annual dues or assessments.
 - b. It may not amend the By Laws.
 - c. It may neither elect new members nor alter the status of existing members, other than to apply the provisions of Article XI.
- 4. The President of the Forum shall serve as Chairman of the Executive Committee and the Secretary of the Forum as its Secretary.

- 5. Meeting of the Executive Committee shall be held at the call of the President of the Forum and each member of the Executive Committee must be notified in writing of the time and place of each such meeting no less than ten (10) days prior to the meeting.
- 6. The annual meeting of the Executive Committee shall precede the annual business meeting of the Forum membership.
- 7. A majority of the voting members of the Executive Committee shall constitute a quorum for the transaction of business.
- 8. The act of a majority of members of the Executive Committee present at a duly called meeting at which a quorum is present shall be the act of the Executive Committee unless the act of a greater number is required by applicable statute or these By Laws.
- 9. Any action which is required by law of the Articles of Incorporation or these By-laws to be taken at a meeting of the Executive Committee, or any other action which may be taken without a meeting if a consent in writing, setting forth the action taken shall be signed by all of the members of the Executive Committee entitled to vote with respect to the subject matter thereof. Any such consent signed by all of the members of the Executive Committee and and constituted meeting of the Executive Committee.
- 10. American Venous Forum Foundation: At its Annual Meeting, the Executive Committee shall elect up to eight (8) individuals to serve as members of the Board of Directors of the American Venous Forum Foundation. These eight individuals shall include the Secretary, Treasurer, and Immediate Past President of the American Venous Forum. Each elected Director, other than the Secretary and Treasurer, shall serve a staggered term of up to three (3) years and shall be eligible for an additional reappointment of one (1) three-year term for a maximum of six (6) years of service to the Board.

ARTICLE VI - COUNCILORS AND OFFICERS

- The officers of the Forum shall be a President, a President Elect, Secretary, Treasurer and Recorder, all to be elected as provided in the By Laws. Said officers shall serve Ex Officio as voting members of the Executive Committee.
- 2. All officers of the Forum, except the Secretary, the Recorder, the Archivist, and the Treasurer, shall be elected for terms of one (1) year each and until their successors are elected and qualified. The President may not serve more than one (1) consecutive term. The Secretary, Recorder and Treasurer will serve three years each and until their successors are elected and qualified. Three Councilors shall be elected serving overlapping terms of three years each.
- 3. A Councilor, Archivist, and the officers of the Forum shall be nominated by the Nominating Committee, which shall present the slate to the Executive Committee at its annual meeting and to the members at the annual business meeting. Additional nominations may be made from the floor at the annual business meeting each year. The election shall take place at the executive session.

Election of officers shall be by a majority of the votes cast. The three candidates for Councilor who receive the most votes shall be elected, provided that each member may vote for three candidates for Councilor and may not cumulate his or her votes.

4. The President shall preside a the meetings of the Forum membership Executive Committee, and Officers, and preserve order, regulate debates, announce results of elections, appoint committees not otherwise provided for in the Bylaws, sign certificates of membership, and perform all other duties normally appertaining to his office.

- 5. The President Elect in the absence or incapacity of the President shall perform the duties of the President's office.
- 6. In the absence of both the President and the President Elect, the position shall be taken by a chairman pro tem, nominated and elected by such members of the Executive Committee as are present.
- 7. The Secretary shall keep the minutes of the meetings of the Forum, the Executive Committee, and the Officers; attest all official acts requiring certification; notify councilors, officers and members of their election and take charge of all papers not otherwise provided for. At least ten (10) days but not more than thirty (30) days prior to each annual or special meeting, the Secretary shall issue to all members of the Society a program of the forthcoming meeting. The Secretary shall compile a written report to be read at the annual business meeting of the Forum in which shall be included the list of candidates proposed for membership, as approved by the Executive Committee.
- 8. The Treasurer shall receive all monies and funds belonging to the Forum to pay all bills; render bills for dues and assessments as soon as possible after the annual meeting; and report to the Executive Committee at each annual meeting the names of all members in arrears as to dues.
- 9. The Recorder shall receive all papers and reports of discussions on paper presented before the Forum or read by title.
- 10. The Archivist shall serve for three years and until a successor is elected and qualified. The Archivist shall be nominated by the Nominating Committee.

ARTICLE VII - COMMITTEES

- The Standing Committees of the Forum shall consist of a Membership Committee, a Nominating Committee, a Program Committee, a Committee on Arrangements of the Annual Meeting, an International Relations Committee, a Committee on Issues, a Committee on Research, and an Honorary Membership Committee.
- The By-Laws Committee shall consist of three members to serve overlapping terms of three years each with the (Secretary of the Forum) serving as Chair. A new member shall be appointed annually by the President. They will review the By-Laws from time to time as directed by the Executive Committee.
- 3. The Membership Committee shall consist of three (3) elected members, who shall serve overlapping terms of three (3) years each, plus the Secretary as an Ex Officio member. The senior member in terms of service on this committee shall be the chairman. The Nominating Committee shall present, annually, one or more candidate(s) to serve as a member of the Membership Committee as part of its slate to the Executive Committee at its annual meeting. Election shall be by the members at large at the executive session. Election shall be by a majority of the votes cast. The functions of the Committee shall be to pass upon the professional and ethical qualifications of the applicants and to advise the Executive Committee of the recommendations of the Committee.
- 4. The Nominating Committee shall consist of the three (3) most recent available Past Presidents and shall be appointed by the President one (1) month before the annual meeting. Its function shall be to make up a slate of officers and a member or members of the Membership Committee to be presented at the annual meeting to the members at the Executive Session. The Senior Member in terms of service on this Committee shall be the Chairman.
- 5. The Program Committee shall consist of three (3) members who shall be appointed, one in each year, by the President to serve overlapping terms of three (3) years each. The

senior member in terms of service on this committee shall be the chairman. The Secretary and Recorder shall be Ex Officio members of the Program Committee. The function of the Program Committee shall be to solicit papers and other presentations from members and other individuals and to make up the program for the annual meeting.

- 6. The Committee on Arrangements for the Annual Meeting shall be appointed by the President and consists of members resident in the general locality in which the annual meeting is to be held, together with President, Secretary and Recorder acting Ex Officio. The function of this committee shall be the making of general arrangements for the annual meeting.
- 7. The International Relations Committee shall consist of at least three (3) members who shall be appointed, one in each year, by the President to serve overlapping terms of three years each. The senior member in terms of service on this committee shall be the chairman. The Secretary of the Forum shall serve as Ex Officio of the Committee. The functions of the International Relations Committee shall be to establish and maintain communications with venous forums and other related vascular organizations outside of the United States for the purposes of the exchange of information.
- 8. The Committee on Issues shall consist of at least four (4) members who shall be appointed, one in each year, by the President to serve overlapping terms of four (4) years each. The senior member in terms of service on this committee shall be the chairman. The Secretary shall serve as an Ex Officio member of this Committee. The Committee on Issues will have, as one of its responsibilities, the monitoring and interpretation of health care related issues. The Committee shall present its observations and recommendations for action to the Executive Committee.
- 9. The Research Committee shall consist of five (5) members who shall be appointed, one in each year, by the President to serve overlapping terms of five (5) years each. The senior member in terms of service on this committee shall be the chairman. The Secretary of the Forum shall serve as an Ex Officio member of this Committee. The responsibilities of this Committee shall be to promote opportunities in research in venous diseases; to define areas of clinical research that require multicenter clinical efforts; and, to promote research investment in venous disease by national granting agencies."
- 10. The Honorary Membership Committee shall consist of the three (3) most immediate Past Presidents on the Executive Committee of the Forum. The most senior member shall serve as Chairman. The Committee shall be responsible for reviewing candidates for Honorary Membership status and recommending actions to the Executive Committee.
- 11. The Executive Committee may from time to time establish such other committees as it deems advisable. Each such committee shall consist of such persons and shall have such duties and powers as may be designated by the Executive Committee upon establishment of the committee or from time to time thereafter. Unless otherwise provided by the Executive Committee, the President shall appoint the members of each such committee.
- 12. Any vacancy occurring among the members of any elected committee of the Forum shall be filled by appointment by the President, the appointee to serve until the next annual meeting of the Forum membership.
- 13. Members of the Executive Committee, Officers or a Committee may participate in any meeting thereof with a conference telephone or similar communications equipment by means of which all persons participating in the meeting can hear each other, and such participation in a Committee meeting shall constitute presence in person at the meeting.

ARTICLE VIII - MEETINGS

- 1. The annual business meeting of the Forum shall be held at a time and place to be determined by the Executive Committee.
- The Executive Committee shall meet in the week prior to the annual meeting, at a time and place designated by the President. The Chairman of the Membership Committee, the Nominating Committee and the Committee on Arrangements shall meet with the Executive Committee in an advisory capacity.
- 3. Twenty five (25) voting members present in person shall constitute a quorum at a meeting of the membership.
- 4. The vote of a majority of members present and voting at a duly called meeting at which a quorum is present shall be necessary for the adoption of any matter voted upon by the members, unless a greater proportion is required by the applicable statute, the Articles of Incorporation, or these Bylaws.
- 5. Members may not cast their votes by proxy.
- 6. The executive session of the Forum shall be held at a time and place to be set by the President. The business of the Forum shall be conducted at this time.
- 7. The scientific sessions at the annual meeting shall consist of presentations of posters and papers and the discussion of these papers. An Active or a Senior member must be a participant, co-author or sponsor of each presentation selected.
- 8. From time to time when deemed advisable by the Executive Committee, eminent investigators in the field of venous disease or allied sciences may be invited to present a special lecture during the annual meeting. This lecture shall be known as the "D. Eugene Strandness, Jr., MD Memorial Lecture. Each speaker who presents such a lecture shall receive an appropriate honorarium and a certificate of appreciation from the Forum.

ARTICLE IX - INVITED GUESTS

- 1. Any member of the Forum may invite one or more guests to attend the annual meeting of the Forum.
- 2. The names of all guests attending the annual meeting shall be entered under a separate heading in the attendance list.
- 3. All invited guests shall be given the privilege of the floor by the President but shall not be present at the annual business meeting.

ARTICLE X - FEES AND DUES

- 1. Initiation fees and assessments shall be proposed by the Executive Committee and approved by the membership at an annual executive session. The Executive Committee shall set dues for membership in all categories from time to time and publish same to the membership at the annual business meeting.
- 2. Any member of the Forum in arrears as to dues for one (1) year shall be notified of that fact by the Treasurer, by registered letter, which shall contain a copy of this Section 2. If the dues are not paid before the next annual business meeting or if some reasonable explanation of the delinquency is not forthcoming, the name of the delinquent member shall be presented at that Executive Committee meeting and, on a majority vote of the Executive Committee, the name may be stricken from the membership list. The Executive Committee

ARTICLE XI - RESIGNATIONS AND DISCIPLINE

- 1. Resignations of members not in arrears as to dues may be accepted at any annual executive committee meeting by a majority vote of the members present.
- 2. Charges of unprofessional or unethical conduct may be brought against any member of the Forum by written complaint signed by three (3) members of the Forum and delivered to the Secretary. The rules governing disciplinary proceedings based upon such charges shall be as established from time to time by the Executive Committee.

ARTICLE XII - PAPERS AND REPORTS

- 1. All papers and reports read before the Forum shall be delivered to the Recorder at the time of their presentations and submitted online as directed by the Recorder.
- 2. No paper shall be published as having been read before the Forum unless it has been read by title or otherwise before the Forum.

ARTICLE XIII - PROCEDURE

The proceedings of the Forum shall be conducted under Robert's Rules of Order Newly Revised and as amended from time to time.

ARTICLE XIV - CERTIFICATE OF MEMBERSHIP

Every elected member of the Forum shall be entitled to a certificate of membership signed by the President and Secretary.

ARTICLE XV - FISCAL YEAR

The fiscal year of this corporation shall begin on the first of January in each year and shall run through the 31st day of December in that year.

ARTICLE XVI - NOTICE AND WAIVER OF NOTICE

- 1. Whenever under applicable law, these By-laws, or a resolution of the Executive Committee, notice is required to be given to any member, Executive Committee member or officer, such notice may be given in writing, by mail, addressed to such member, Executive Committee member or officer at his or her address as it appears on the records of the Forum. Such mailed notice shall be deemed to have been given when deposited in the United States mail in a sealed envelope so addressed, with postage thereon prepaid.
- 2. Whenever, under applicable law, these By-laws or a resolution of the Executive Committee, any notice is required to be given, a waiver thereof in writing, signed by the person or persons entitled to such notice, whether before or after the time stated therein, shall be deemed equivalent to the giving of such notice. In addition, the attendance of a member or Executive Committee member at any meeting shall constitute a waiver of notice of such meeting, except where an individual attends the meeting for the express purpose of objecting to the transaction of any business because the meeting is not lawfully called or convened.

ARTICLE XVII - INDEMNIFICATION

- 1. To the full extent specifically authorized by, and in accordance with the procedures prescribed in Section 108.75 of the Illinois General Not for Profit Corporation Act of 1986 (or the corresponding provisions of any future statute applicable to corporations organized under the Act), the Forum shall indemnify any and all members of the Executive Committee (which members shall hereinafter in this Article be referred to as "Directors") and any and all of its officers, committee members, employees, agents and other authorized representatives for expenses and other amounts paid in connection with legal proceedings (whether threatened, pending or completed) in which any such person became involved by reason of serving in any such capacity for the Forum.
- 2. Upon specific authorization by the Executive Committee, the Forum may purchase and maintain insurance on behalf of any or all directors, officers, employees, agents or representatives of the Forum against any liability asserted against any such person and incurred in any such capacity, or arising out of the status of serving in any such capacity, whether or not the Forum would have the power to indemnify them against such liability under the provisions of Section I of this Article.

ARTICLE XVIII - AMENDMENT

These By-laws may be amended by a three-fourths vote of the members present and voting at a properly called and convened of an annual business meeting or special meeting of the Forum provided that the proposed amendment has been submitted to the Secretary by at least three (3) voting members of the Forum at least three (3) months prior to the executive session of the Forum. The Secretary shall mail the proposed amendment to all voting members at least thirty (30) days prior to the executive session, accompanied by notice that such amendment will be acted upon at that **Annual Business Meeting**.

THE AMERICAN VENOUS FORUM

PROVISO TO THE BY-LAWS

ARTICLE I - EFFECT OF PROVISO

This Proviso to the By-laws (the "By-laws") of the American Venous Forum, an Illinois not for profit corporation (the "Forum"), shall control and supersede the rules and regulations for the governance of the Forum contained in the By-laws as of the date on which they are adopted. Except as specifically modified by this Proviso, all other provisions of the By-laws shall remain in full force and effect.

ARTICLE II - OFFICERS

The initial members of the Executive Committee of the Forum, which members are named in the Articles of Incorporation of the Forum as filed with the Illinois Secretary of State on February 7, 1989 shall elect the initial officers of the Forum from among the members of the Executive Committee. The officers so elected shall serve until the next annual executive session of the members of the Forum and until their successors shall have been elected and qualified.

DRAFTED: October 23, 1988 ADOPTED: February 22, 1989 AMENDED: February 19, 1999 AMENDED: February 16, 2007

AMERICAN VENOUS FORUM

20th ANNUAL MEETING February 20-23, 2008

AVF EXHIBITING COMPANIES

Bacchus Vascular, Inc. was founded by Dr. Thomas Fogarty to transform blood clot removal from peripheral blood vessels into a fast, simple procedure. Thousands of patients suffering from DVT have been treated using the Trellis® Peripheral Infusion System, a catheter-based device designed for single setting treatment.

BiaCare Medical is a leading manufacturer of custom and standard compression wear, including MedAssist, CompreFit, CompreFit and FoamSleeve brands. BiaCare Medical also produces compression therapy supplies, including LympheBand, LympheSoft, and Silvernette products. BiaCare Medical products are available through leading compression dealers. www.BiaCare.com

Jobst is the worldwide leader in venous and lymphatic health products. Jobst has combined innovation and quality to offer a complete range of gradient compression products, providing solutions for patients with various vascular disorders. When you think of comfort, health and style – think of JOBST!

43 Carolon Company is the compression hosiery manufacturer of choice and the only U.S. owned manufacturer of compression hosiery. Health Support hosiery is easier to get on, more comfortable to wear, washer and dryer safe and much less expensive.
CIRCAID MEDICAL PRODUCTS
44 Cook® Medical has the complete line of interventional products from stick to stent. Cook® Medical offers access products like needles, Micropuncture, wire guides, catheters, dilators, introducers and sheaths. Our therapeutic devices include Zilver and Formula stents, Zenith stent-graft, Advance PTA balloon, Tulip IVC Filter, and a complete line of embolization coils.
COVIDIEN, FORMERLY TYCO HEALTHCARE/ KENDALL
DIOMED INC

GE HEALTHCARE
HEALTHPOINT, LTD
HK SURGICAL

HUNTLEIGH HEALTHCARE, INC......4

Laser Peripherals designs, manufactures, distributes and OEM's medical laser fibers for use in both hard and soft tissue surgical laser applications. We manufacture and market at least thirty different surgical fiber options for use with Diode, Holmium, KTP and Nd:YAG lasers. Fiber designs include freebeam, contact or lateral emitting.

Lympha Press USA manufactures and distributes Lympha Press compression therapy devices for treatment of lymphedema and venous disorders. Our systems are renowned worldwide for quality and efficacy, and are reimbursed by Medicare and private insurance. See The Petite Basic System, our affordable home care device; Lympha Press Mini, our 12-chamber calibrated gradient system; and Lympha Press Plus, a programmable system for clinic use, as well as unique garments you won't find anywhere else.

Medi is the worldwide leader in medical compression therapy and prevention of venous ulcers. We are dedicated to innovative Phlebology and Lymphology product development, conscientious quality control and total customer satisfaction. When asked how physicians can provide therapeutic compression in product patients will actually wear. Physicians answer with Medi.

Apligraf® is the first and only fully differentiated living bi-layered cell therapy that is FDA approved for the treatment of venous leg ulcers and diabetic foot ulcers. Apligraf is easy to incorporate into practice. Positive reimbursement in all settings. Well tolerated in over 150,000 patient applications. www.organogenesis.com www.apligraf.com

Possis Medical offers the widest range of indications and therapies for thrombus, with products like the Fetch™ Aspiration Catheter, PowerPulse® Delivery, GuardDOG® Occlusion System, SafeSeal™ Hemostasis dressing, and the new, easy-to-use AngioJet® Ultra Rheolytic™ Thrombectomy System with seven specialized catheters providing choice in length, platform, clot removal power and profile.

OFI-AVENTIS	•

SONOSITE	33
----------	----

OMBOSISCLINIC.COM

 TOTAL VEIN SOLUTIONS
 11

 Total Vein Systems offers the most extensive line of Custom and Universal Procedure Packs for Endovascular Surgery and other procedures associated with Varicose Vein Disease.

 Total Vein also markets a complete line of premium laser fibers, access devices and venous surgical supplies.

Vascular Solutions offers the most comprehensive line of endovenous laser therapy products available, from the Vari-Lase® Console to procedure kits and accessories like the Klein Pump. We also support customers with a free Vein Practice Business Plan, reimbursement newsletter, Vein Care Essentials training, patient education, marketing materials and website patient referrals.

VNUS® Medical Technologies is the proven leader in minimally-invasive treatment of venous reflux. The VNUS Closure(r) System combines a proprietary radiofrequency (RF) generator with a family of disposable catheters to close diseased veins using temperature-controlled RF energy. Physicians can treat incompetent saphenous, perforator and tributary vessels with minimal pain and bruising.

VOLCANO THERAPEUTICS, INC
Founded in 2000, Volcano Corporation (NASDAQ:VOLC) develops, manufactures and commercializes a broad suite of intravascular ultrasound and functional measurement products that enhance the diagnosis and treatment of vascular and structural heart disease. For more information, visit the company's website at www.volcanocorp.com
WAGNER MEDICAL
We specialize in vein-sclerotherapy products, such as the latest textbooks, bi-directional and unidirectional dopplers, transducers, PPG-LRR detection of acute DVT and venous insufficiency and reconditioned duplex ultrasound. Vein hooks for ambulatory phlebectomies. www.wagnermedical.net
WELLS JOHNSON COMPANY

Since 1983, Wells Johnson Company has sold over 9400 Aspirators World Wide, and is dedicated to deliver quality, reliability and great customer service. Wells Johnson Company sells the finest in Aspirators, Cannulas, Infusion Pumps, Garments and Disposables.

AMERICAN VENOUS FORUM

Authors Index

Abdul Rahman, M. N	
Abdullah, N. M	8
Adams, E.	23
Adams, G.	2
Adams, J.	24
Aidinian, G.	23
Allaert, F. A.	P-10
Allan, P.	34
Alm, J	
Almeida, J. I.	P-26
Amin, A. A.	23
Amsler, F	
Andrews, P. C	8
Aonuma, K	
Araujo, S. P.	
Aravind, B	20
Ascher, E	3 <i>,</i> P-31
Assi, Z. I	
Azuma, N	
Ballard, N. E.	P-1
Beidler, S.	9
Benigni, J	P-17
Bergan, J.	P-28
Berges, D	24
Berndt, D.	9
Blackburn, S.	2
Blaettler, W	P-20
Bloom, J.	
Bogdanovic, D. C.	35
Bowser, A. N.	P-30
Brown, S.	P-6
Burnand, K.	10
Burseta, P	P-10
Bush, R.	
Capella, A	
Caprini, J. A.	
Carmichael, S	34

Guerzoni, S.	
Gulati, S. 5, P-5 Hamilton, R. 16 Harmond, K. 14 Han, G. P-8 Hartung, O. 30 Hatfield, J. 5, P-5 Hawley, A. E. 8, 11, P-1 Henke, P. K. 8, 11, P-1, P-6 Hertzman, P. A. 21, P-27 Hill, D. A. 16 Hingorani, A. 3, P-31 Homma, S. 28 Hong, M. 2 Hoshino, S. P-25 Howard, A. 24 Hurd, A. P-29 Ichiki, M. P-25 Isaka, Y. 28 Jacobson, C. P-8 Jankovic, R. P-14 Johnson, C. P-30 Jones, L. 33 Jones, W. T. P-21 Jovanovic, M. M. P-14 Kachman, M. 8 Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. 32 Khalil, R. 12 Kistner, R. L. 17 Klem, T. M. 15 <td></td>	
Hamilton, R. 16 Harn, G. P-8 Hartung, O. 30 Hatfield, J. 5, P-5 Hawley, A. E. 8, 11, P-1 Henke, P. K. 8, 11, P-1, P-6 Hertzman, P. A. 21, P-27 Hill, D. A. 16 Hingorani, A. 3, P-31 Homma, S. 28 Hong, M. 2 Hoshino, S. P-25 Howard, A. 24 Hurd, A. P-29 Ichiki, M. P-25 Isaka, Y. 28 Jacobson, C. P-8 Jankovic, R. P-14 Johnson, C. P-30 Jones, L. 33 Jones, W. T. P-21 Jovanovic, M. M. P-14 Kachman, M. 8 Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. 32 Khalil, R. 12 Kistner, R. L. 17 Klem, T. M. 15 Knipp, B. S. 2 Kono, T. 29, P-4, P-	Guex, J. J4
Hammond, K. 14 Han, G. P-8 Hartung, O. 30 Hatfield, J. 5, P-5 Hawley, A. E. 8, 11, P-1 Henke, P. K. 8, 11, P-1 Henke, P. K. 8, 11, P-1, P-6 Hertzman, P. A. 21, P-27 Hill, D. A. 16 Hingorani, A. 3, P-31 Homma, S. 28 Hong, M. 2 Hoshino, S. P-25 Howard, A. 24 Hurd, A. P-29 Ichiki, M. P-25 Isaka, Y. 28 Jacobson, C. P-8 Jankovic, R. P-14 Johnson, C. P-30 Jones, L. 33 Jones, W. T. P-21 Jovanovic, M. M. P-14 Kachman, M. 8 Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. 32 Khalil, R. 12 Kistner, R. L. 17 Klem, T. M. 15 Knipp, B. S.	Gulati, S 5, P-5
Han, G. P-8 Hartung, O. 30 Hatfield, J. 5, P-5 Hawley, A. E. 8, 11, P-1 Henke, P. K. 8, 11, P-1 Henke, P. K. 8, 11, P-1 Henke, P. K. 16 Hingorani, A. 3, P-31 Homma, S. 28 Hong, M. 2 Hoshino, S. P-25 Howard, A. 24 Hurd, A. P-29 Ichiki, M. P-25 Isaka, Y. 28 Jacobson, C. P-8 Jankovic, R. P-14 Johnson, C. P-30 Jones, L. 33 Jones, W. T. P-21 Jovanovic, M. M. P-14 Kachman, M. 8 Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. 32 Khalil, R. 12 Kistner, R. L. 17 Klem, T. M. 15 Knipp, B. S. 2 Kono, T. P-9 Lane, C. P-6	Hamilton, R
Hartung, O. .30 Hatfield, J. .5, P-5 Hawley, A. E. .8, 11, P-1 Henke, P. K. .11, P-1, P-6 Hertzman, P. A. .21, P-27 Hill, D. A. .16 Hingorani, A. .3, P-31 Homma, S. .28 Hong, M. .2 Hoshino, S. P-25 Howard, A. .24 Hurd, A. .29 Ichiki, M. P-25 Isaka, Y. .28 Jacobson, C. .P-8 Jankovic, R. .9-14 Johnson, C. .93 Jones, L. .33 Jones, W. T. .9-21 Jovanovic, M. M. .9-14 Kachman, M. .8 Kakkos, S. K. .9-7 Kaspar, S. .9-16 Kazanjian, S. .32 Khalil, R. .12 Kistner, R. L. .17 Klem, T. M. .15 Knipp, B. S. .2 Kono, T. .92, P-4, P-11 Kurtoglu, M. .9-9 Lan	Hammond, K14
Hatfield, J. 5, P-5 Hawley, A. E. 8, 11, P-1 Henke, P. K. 8, 11, P-1 Henke, P. K. 11, P-1 Hertzman, P. A. 21, P-27 Hill, D. A. 16 Hingorani, A. 3, P-31 Homma, S. 28 Hong, M. 2 Hoshino, S. P-25 Howard, A. 24 Hurd, A. P-29 Ichiki, M. P-25 Isaka, Y. 28 Jacobson, C. P-8 Jankovic, R. P-14 Johnson, C. P-30 Jones, L. 33 Jones, L. 33 Jones, W. T. P-21 Jovanovic, M. M. P-14 Kachman, M. 8 Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. 32 Khalil, R. 12 Kistner, R. L. 17 Klem, T. M. 15 Knipp, B. S. 2 Kono, T. P-9 Lane, C. P-6	Han, G
Hawley, A. E. 8, 11, P-1 Henke, P. K. 8, 11, P-1, P-6 Hertzman, P. A. 21, P-27 Hill, D. A. 16 Hingorani, A. 3, P-31 Homma, S. 28 Hong, M. 2 Hoshino, S. P-25 Howard, A. 24 Hurd, A. P-29 Ichiki, M. P-25 Isaka, Y. 28 Jacobson, C. P-8 Jankovic, R. P-14 Johnson, C. P-30 Jones, L. .33 Jones, W. T. P-21 Jovanovic, M. M. P-14 Kachman, M. 8 Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. .32 Khalil, R. .12 Kistner, R. L. .17 Klem, T. M. .15 Knipp, B. S. .2 Kono, T. .29, P-4, P-11 Kurtoglu, M. P-9 Lane, C. P-6 Lane, R. J. P-22 Lebard, C.	Hartung, O
Henke, P. K. 8, 11, P-1, P-6 Hertzman, P. A. 21, P-27 Hill, D. A. 16 Hingorani, A. 3, P-31 Homma, S. 28 Hong, M. 2 Hoshino, S. P-25 Howard, A. 24 Hurd, A. P-29 Ichiki, M. P-25 Isaka, Y. 28 Jacobson, C. P-8 Jankovic, R. P-14 Johnson, C. P-30 Jones, L. 33 Jones, W. T. P-21 Jovanovic, M. M. P-14 Kachman, M. 8 Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. 32 Khalil, R. 12 Kistner, R. L. 17 Klem, T. M. 15 Knipp, B. S. 2 Kono, T. 29, P-4, P-11 Kurtoglu, M. P-9 Lane, C. P-6 Laredo, J. P-22 Lebard, C. P-18 LeBaron, S. P-18	Hatfield, J 5, P-5
Henke, P. K. 8, 11, P-1, P-6 Hertzman, P. A. 21, P-27 Hill, D. A. 16 Hingorani, A. 3, P-31 Homma, S. 28 Hong, M. 2 Hoshino, S. P-25 Howard, A. 24 Hurd, A. P-29 Ichiki, M. P-25 Isaka, Y. 28 Jacobson, C. P-8 Jankovic, R. P-14 Johnson, C. P-30 Jones, L. 33 Jones, W. T. P-21 Jovanovic, M. M. P-14 Kachman, M. 8 Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. 32 Khalil, R. 12 Kistner, R. L. 17 Klem, T. M. 15 Knipp, B. S. 2 Kono, T. 29, P-4, P-11 Kurtoglu, M. P-9 Lane, C. P-6 Laredo, J. P-22 Lebard, C. P-18 LeBaron, S. P-18	Hawley, A. E
Hill, D. A. 16 Hingorani, A. 3, P-31 Homma, S. 28 Hong, M. 2 Hoshino, S. P-25 Howard, A. 24 Hurd, A. P-29 Ichiki, M. P-25 Isaka, Y. 28 Jacobson, C. P-8 Jankovic, R. P-14 Johnson, C. P-30 Jones, L. 33 Jones, W. T. P-21 Jovanovic, M. M. P-14 Kachman, M. 8 Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. 32 Khalil, R. 12 Kistner, R. L. 17 Klem, T. M. 15 Knipp, B. S. 2 Kono, T. 29, P-4, P-11 Kurtoglu, M. P-9 Lane, C. P-6 Laredo, J. P-22 Lebard, C. P-18 LeBaron, S. P-18	
Hill, D. A. 16 Hingorani, A. 3, P-31 Homma, S. 28 Hong, M. 2 Hoshino, S. P-25 Howard, A. 24 Hurd, A. P-29 Ichiki, M. P-25 Isaka, Y. 28 Jacobson, C. P-8 Jankovic, R. P-14 Johnson, C. P-30 Jones, L. 33 Jones, W. T. P-21 Jovanovic, M. M. P-14 Kachman, M. 8 Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. 32 Khalil, R. 12 Kistner, R. L. 17 Klem, T. M. 15 Knipp, B. S. 2 Kono, T. 29, P-4, P-11 Kurtoglu, M. P-9 Lane, C. P-6 Laredo, J. P-22 Lebard, C. P-18 LeBaron, S. P-18	Hertzman, P. A
Hingorani, A. 3, P-31 Homma, S. 28 Hong, M. 2 Hoshino, S. P-25 Howard, A. 24 Hurd, A. P-29 Ichiki, M. P-25 Isaka, Y. 28 Jacobson, C. P-8 Jankovic, R. P-14 Johnson, C. P-30 Jones, L. 33 Jones, V. T. P-21 Jovanovic, M. M. P-14 Kachman, M. 8 Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. 32 Khalil, R. 12 Kistner, R. L. 17 Klem, T. M. 15 Knipp, B. S. 2 Kono, T. 29, P-4, P-11 Kurtoglu, M. P-9 Lane, C. P-6 Laredo, J. P-22 Lebard, C. P-18 LeBaron, S. P-6	
Homma, S.	
Hoshino, S. P-25 Howard, A. .24 Hurd, A. P-29 Ichiki, M. P-25 Isaka, Y. .28 Jacobson, C. P-8 Jankovic, R. P-14 Johnson, C. P-30 Jones, L. .33 Jones, W. T. P-21 Jovanovic, M. M. P-14 Kachman, M. .8 Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. .32 Khalil, R. .12 Kistner, R. L. .17 Klem, T. M. .15 Knipp, B. S. .2 Kono, T. .29, P-4, P-11 Kurtoglu, M. P-9 Lane, C. P-6 Laredo, J. P-22 Lebard, C. P-18 LeBaron, S. P-6	
Hoshino, S. P-25 Howard, A. .24 Hurd, A. P-29 Ichiki, M. P-25 Isaka, Y. .28 Jacobson, C. P-8 Jankovic, R. P-14 Johnson, C. P-30 Jones, L. .33 Jones, W. T. P-21 Jovanovic, M. M. P-14 Kachman, M. .8 Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. .32 Khalil, R. .12 Kistner, R. L. .17 Klem, T. M. .15 Knipp, B. S. .2 Kono, T. .29, P-4, P-11 Kurtoglu, M. P-9 Lane, C. P-6 Laredo, J. P-22 Lebard, C. P-18 LeBaron, S. P-6	Hong, M2
Hurd, A. P-29 Ichiki, M. P-25 Isaka, Y. 28 Jacobson, C. P-8 Jankovic, R. P-14 Johnson, C. P-30 Jones, L. 33 Jones, W. T. P-21 Jovanovic, M. M. P-14 Kachman, M. 8 Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. 32 Khalil, R. 12 Kistner, R. L. 17 Klem, T. M. 15 Knipp, B. S. 2 Kono, T. 29, P-4, P-11 Kurtoglu, M. P-9 Lane, C. P-6 Lane, R. J. P-2 Lebard, C. P-18 LeBaron, S. P-6	
Ichiki, M. P-25 Isaka, Y. 28 Jacobson, C. P-8 Jankovic, R. P-14 Johnson, C. P-30 Jones, L. 33 Jones, V. T. P-21 Jovanovic, M. M. P-14 Kachman, M. 8 Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. 32 Khalil, R. 12 Kistner, R. L. 17 Klem, T. M. 15 Knipp, B. S. 2 Kono, T. 29, P-4, P-11 Kurtoglu, M. P-9 Lane, C. P-6 Lane, R. J. P-2 Lebard, C. P-18 LeBaron, S. P-6	
Ichiki, M. P-25 Isaka, Y. 28 Jacobson, C. P-8 Jankovic, R. P-14 Johnson, C. P-30 Jones, L. 33 Jones, V. T. P-21 Jovanovic, M. M. P-14 Kachman, M. 8 Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. 32 Khalil, R. 12 Kistner, R. L. 17 Klem, T. M. 15 Knipp, B. S. 2 Kono, T. 29, P-4, P-11 Kurtoglu, M. P-9 Lane, C. P-6 Lane, R. J. P-2 Lebard, C. P-18 LeBaron, S. P-6	Hurd, AP-29
Isaka, Y.	
Jankovic, R	Isaka, Y28
Johnson, C	Jacobson, C P-8
Jones, L	Jankovic, R P-14
Jones, W. T	Johnson, C P-30
Jovanovic, M. M	Jones, L
Kachman, M.	Jones, W. T P-21
Kakkos, S. K. P-7 Kaspar, S. P-16 Kazanjian, S. 32 Khalil, R. 12 Kistner, R. L. 17 Klem, T. M. 15 Knipp, B. S. 2 Kono, T. 29, P-4, P-11 Kurtoglu, M. P-9 Lane, C. P-6 Lanedo, J. P-22 Lebard, C. P-18 LeBaron, S. P-6	Jovanovic, M. MP-14
Kaspar, S. P-16 Kazanjian, S. 32 Khalil, R. 12 Kistner, R. L. 17 Klem, T. M. 15 Knipp, B. S. 2 Kono, T. 29, P-4, P-11 Kurtoglu, M. P-9 Lane, C. P-6 Lanedo, J. P-22 Lebard, C. P-18 LeBaron, S. P-6	Kachman, M8
Kazanjian, S.	
Khalil, R. 12 Kistner, R. L. 17 Klem, T. M. 15 Knipp, B. S. 2 Kono, T. 29, P-4, P-11 Kurtoglu, M. P-9 Lane, C. P-6 Lane, R. J. P-2, P-3 Laredo, J. P-22 Lebard, C. P-18 LeBaron, S. P-6	Kaspar, SP-16
Kistner, R. L. 17 Klem, T. M. 15 Knipp, B. S. 2 Kono, T. 29, P-4, P-11 Kurtoglu, M. P-9 Lane, C. P-6 Lanedo, J. P-22 Lebard, C. P-18 LeBaron, S. P-6	Kazanjian, S
Klem, T. M. 15 Knipp, B. S. 2 Kono, T. 29, P-4, P-11 Kurtoglu, M. P-9 Lane, C. P-6 Lane, R. J. P-2, P-3 Laredo, J. P-22 Lebard, C. P-18 LeBaron, S. P-6	Khalil, R12
Knipp, B. S.	
Kono, T	
Kurtoglu, M. P-9 Lane, C. P-6 Lane, R. J. P-2, P-3 Laredo, J. P-22 Lebard, C. P-18 LeBaron, S. P-6	
Lane, C	Kono, T
Lane, R. J	
Laredo, J	
Lebard, C	
LeBaron, S P-6	
Lebow, M 13, P-29	Lebow, M 13, P-29

Lee, A	
Lee, B.	P-22
Lee, J	31, P-8
Lees, T.	1
Li, X	P-31
Liboni, A	
Lugli, M.	36, P-10
Lurie, F	
Makimura, S	P-25
Maksimovic, Z. D	
Maleti, O	
Mansoor, F.	2
Marks, N	
Marston, W.	9
Martinez, J.	
Mattaliano, V.	P-13
McCollum, P. T.	5, P-5
McDonald, A. P	P-1
McKinstry, B.	34
Meier, T. R	11
Mekako, A. I.	
Menegatti, E	7
Milic, D. J.	35, P-14
Milojkovic, V. D	
Miranda Jr, F	18
Mitake, T	28
Monaco, C	
Morrison, N.	
Mosti, G.	
Myers, D. D	
Navin, T	
Neglén, P	
Neville, R	
Nicolaides, A. N.	
Nighswander, C.	
Nozaki, M	
Ogawa, T	
Oh, H	
Osaka, T	
Palazzo, A	
Paleolog, E.	
Paolini, D.	

Partsch, H P-13, P-23
Passman, M19
Patel, N P-31
Pejic, M. A35
Perisic, Z. D P-14
Perrin, M P-10
Peterson, B21
Pfeifer, J
Phillips, M. NP-3
Pichot, O P-18
Pittaluga, P4
Popvic, V. M
Proebstle, T. M P-18
Puggioni, A
Raffetto, J. D
Raines, J. K
Raju, S
Rasmussen, T. E
Reddy, D. J P-7
Reynolds, G P-6
Ricci, M. A
Rich, P
Righi, E
Robertson, L
Ruckely, C
Sakurai, H
Salles-Cunha, S. X. P-32
Sasajima, TP-25
Schnater, J. M
Schutte, P. R
Scott, V
Scriver, G
Sepanski, D. M
Shackford, S
Shiferson, A
Shigematsu, H
Shiina, T
Shakaku, S
Siller, J. P-16
Sivrikoz, E
Slusarczyk, M24
Smith, A 10

Soejima, K	
Stamenkovic, D. M	P-14
Stanley, A	
Stansby, G. P	P-7
Steinthorsson, G	24
Stevens, S	13, P-29
Strahler, J. R	8
Subramonia, S	1
Suda, M	
Sugawara, H	P-25
Tacconi, G	7
Tackett, P	25, 26
Takeuchi, M	29, P-4, P-11
Tan, J	10
Thibert, J. N	11
Tonomura, A	
Tran, V	P-31
Uhl, J	P-17
Uno, K	
Vago, B	
van der Ham, A. C	15
Vandy, F. C	
Vanscheidt, W	
Wakefield, T. W2	, 8, 11, P-1, P-6
Walker, A	8
White, P. W	23
Williams, D	
Wittens, C. H. A	
Wooster, D. L.	22
Wrobleski, S. K	8, 11, P-1
Yamakawa, M	
Yamaki, T	29, P-4, P-11
Yoon, H	17
Zamboni, P	
Zierler, B	
Zivic, S. S	35, P-14

AMERICAN VENOUS FORUM

AUTHOR DISCLOSURES

Abst.#	Author	Disclosure
1	S. Subramonia T. Lees	Stock Options: VNUS Medical Stock Options: VNUS Medical
2	B.S Knipp F. Mansoor M. Hong J. Bloom E. Fellows S. Blackburn G. Adams J. Pfeifer D. Williams T. Wakefield	Nothing To Disclose. Nothing To Disclose.
3	A. Puggioni N. Marks A. Hingorani A. Shiferson E. Ascher	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
4	P. Pittaluga S. Chastanet J. J. Guex	Nothing To Disclose.
5	A. I. Mekako J. Hatfield M. N. Abdul Rahman S. Gulati P. T. McCollum I. C. Chetter	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
6	V. Cheng	Nothing To Disclose.
7	S. Gianesini G. Tacconi A. Palazzo P. Fortini E. Righi E. Menegatti A. Liboni P. Zamboni	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.

Abst.#	Author	Disclosure
8	N. M. Abdullah M. Kachman A. Walker A. E. Hawley S. K. Wrobleski D. D. Myers, Jr. J. R. Strahler P. C. Andrews P. K. Henke T. W. Wakefield	Nothing To Disclose. Nothing To Disclose.
9	S. Beidler C. Douillet D. Berndt P. Rich W. Marston	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
10	J. Tan A. Smith K. Burnand	Nothing To Disclose.
11	A. P. McDonald T. R. Meier A. E. Hawley J. N. Thibert D. M. Farris S. K. Wrobleski P. K. Henke T. W. Wakefield D. D. Myers, Jr.	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
12	J. D. Raffetto R. Khalil	
13	M. Lebow D. Cassada O. Grandas S. Stevens M. Freeman M. Goldman	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
14	R. Bush K. Hammond	
15	T. M. A. L. Klem J. M. Schnater P. R. Schutte A. C. van der Ham C. H. A. Wittens.	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
16	D. A. Hill R. Hamilton	

Abst.#	Author	Disclosure
17	F. Lurie H. Yoon V. Scott R. L. Kistner	Nothing To Disclose. Nothing To Disclose.
18	M. Figueiredo S. P. Araujo F. Miranda, Jr	Nothing To Disclose.
19	M. Passman	Nothing To Disclose.
20	B. Aravind T. Navin C. Monaco E. Paleolog A. H. Davies	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
21	P. A. Hertzman B. Peterson	Nothing To Disclose. Nothing To Disclose.
22	D. L. Wooster	Nothing To Disclose.
23	G. Aidinian A. A. Amin P. W. White E. Adams C. J. Fox M. Cox D. L. Gillespie	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
24	G. Scriver A. Stanley M. A. Ricci K. Corrow M. Slusarczyk S. Shackford J. Adams G. Steinthorsson D. Berges A. Howard	Nothing To Disclose. Nothing To Disclose.
25	P. Neglen P. Tackett S. Raju	Nothing To Disclose.
26	S. Raju P. Tackett P. Neglen	Nothing To Disclose.
27	Withdrawn	N/A

Abst.#	Author	Disclosure
28	T. Osaka	Employment: Hitachi Medical Corp. Employment: Hitachi Medical Corp. Employment: Hitachi Medical Corp. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
29	T. Yamaki M. Nozaki H. Sakurai M. Takeuchi K. Soejima T. Kono	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
30	O. Hartung	Nothing To Disclose.
31	J. Lee B. Zierler	
32	J. Martinez A. J. Comerota S. Kazanjian R. DiSalle D. M. Sepanski Z. I. Assi	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
33	D. Paolini L. Jones A. J. Comerota	Nothing To Disclose.
34	L. Robertson A. Lee K. Gallagher S. Carmichael C. Evans B. McKinstry S. Fraser P. Allan C. Ruckley F. Fowkes	Nothing To Disclose. Nothing To Disclose.
35	D. J. Milic S. S. Zivic D. C. Bogdanovic V. D. Milojkovic M.A. Pejic V. M. Popvic	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.

Abst.#	Author	Disclosure
36	M. Lugli S. Guerzoni O. Maleti	Nothing To Disclose.
P-1	A. E. Hawley D. M. Farris N. E. Ballard A. P. McDonald S. K. Wrobleski P. K. Henke D. D. Myers T. W. Wakefield	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
P-2	R. J. Lane M. L. Cuzilla	
P-3		Ownership/Partnership: AllVascular Pty Ltd Employment: AllVascular Pty Ltd Nothing To Disclose.
P-4	T. Yamaki M. Nozaki H. Sakurai M. Takeuchi K. Soejima T. Kono	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
P-5	A. Mekako J. Hatfield S. Gulati M. Abdul Rahman P. T. McCollum I. C. Chetter	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
P-6	J. Bloom F. C. Vandy S. Brown A. Clay C. Lane G. Reynolds S. LeBaron C. Nighswander P. K. Henke T. W. Wakefield	Nothing To Disclose. Nothing To Disclose.
P-7	S. K. Kakkos J. A. Caprini G. Geroulakos A. N. Nicolaides G. P. Stansby D. J. Reddy	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.

Abst.#	Author	Disclosure
P-8	B. K. Zierler J. Lee G. Han H. Oh C. Jacobson	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
P-9	E. Sivrikoz M. Kurtoglu	
P-10	M. Cazaubon M. Lugli P. Burseta M. Perrin F. A. Allaert	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
P-11	T. Yamaki M. Nozaki H. Sakurai M. Takeuchi K. Soejima T. Kono	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
P-12	G. Mosti	Nothing To Disclose.
P-13	G. Mosti H. Partsch V. Mattaliano	Nothing To Disclose.
P-14	D. J. Milic S. S. Zivic Z. D. Perisic R. Jankovic G. Djordjevic D. M. Stamenkovic Z. D. Maksimovic	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
P-15	Withdrawn	N/A
P-16	S. Kaspar J. Siller Z. Cervinkova	Nothing To Disclose.
P-17	J. Benigni J. Uhl J. Gobin A. Capella	Nothing To Disclose. Nothing To Disclose.
P-18	B. Vago J. Alm O. Goeckeritz C. Lebard	Research Grant & Honoraria: VNUS Medical Nothing To Disclose. Research Grant: VNUS Medical Research Grant: VNUS Medical Research Grant: VNUS Medical Research Grant: VNUS Medical

Abst.#	Author	Disclosure
P-19	J. T. Christenson	Nothing To Disclose.
P-20	W. Blaettler Management H. E. Gerlach	Consulting Fee & Honoraria: Ganzoni
	F. Amsler	
P-21	W. T. Jones M. A. Ricci W. D. Clouse T. E. Rasmussen	Nothing To Disclose. Nothing To Disclose.
P-22	B. Lee J. Laredo D. Deaton R. Neville	Nothing To Disclose. Nothing To Disclose.
P-23	H. Partsch W. Vanscheidt	
P-24	S. Shokoku	Nothing To Disclose.
P-25	T. Ogawa S. Hoshino S. Makimura H. Shigematsu N. Azuma T. Sasajima H. Sugawara M. Ichiki S. Shokoku	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
P-26	J. I. Almeida J. K. Raines	Research Grants: AngioDynamics Research Grants: AngioDynamics
P-27	P. A. Hertzman	Nothing To Disclose.
P-28	V. Cheng J. Bergan	
P-29	M. Lebow A. Hurd D. Cassada M. Freeman O. Grandas S. Stevens M. Goldman	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
P-30	S. Chen A. N. Bowser W. D. Clouse C. Johnson T. E. Rasmussen	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.

Abst.#	Author	Disclosure
P-31	N. Patel A. Puggioni X. Li A. Hingorani A. Shiferson V. Tran E. Ascher	Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose. Nothing To Disclose.
P-32	S. X. Salles-Cunha N. Morrison	

NOTES



IS YOUR AVF MEMBERSHIP INFORMATION CURRENT?

For Example:

- Do you have a new email address?
- Do you have a new address or phone number?

Please let us know so that your AVF records stay current, and that all important updates and news reach you!

PLEASE PRINT				
First	М.		Last	Suffix
Email Address				
Daytime Phone		Fax		
MAILING ADDRESS				
Institution				
Street				
City	State		Zip	Country

Please return your completed form to the AVF Registration Desk, or fax your form to 978-744-5029.

NOTES