# South Carolina Bleeding Disorders 2021 Medical Symposium

Understanding Hemophilia and Inherited Bleeding Disorders: the Treatment, the South Carolina Comprehensive Care Teams. and Potential Complications





Free!

Save The Date

PHILIA Friday April 23, 4:00pm - 5:30pm

Carolina Saturday April 24, 9:00am - 12:00pm

This medical symposium is an opportunity to bring hospital administration, healthcare professionals, ER physicians, hematologists, and primary care physicians practicing in South Carolina who currently have patients in their practice or emergency care, or potential future patients who have a diagnosis of hemophilia or related inherited bleeding disorders specialized medical education in the healthcare needs of this patient population.

(Continuing Medical Education Credits (1unit) will be provided)

### Friday Panelist







Including Robin Jones, MSN, MHA, RN, CPN, CPHON - SC Hemophilia Treatment Center Nurse Coordinator

The Comprehensive Model of Care for Hemophilia and Bleeding Disorders: The Hemophilia Treatment Centers in South Carolina

Meet the South Carolina Hemophilia Treatment Centers and the healthcare team who treat patients who have a diagnosis of Hemophilia, von Willebrand Disease, platelet disorders, and other rare factor deficiencies.

## Saturday Speakers



University of Miami, HTC

9:00am: The Hemophilia and Rare Bleeding Disorders Awareness Series

(1 CU Credit)





Persons with Bleeding Disorders

Emergency and Orthopaedic Issues for

11:00am:

10:00am: Shemophilia: Women with Bleeding Disorders

# Symposium Schedule

#### Friday:

4:00pm: Welcome & Introductions: Hemophilia of South Carolina and the SC Advocacy Coalition

#### The Comprehensive Model of Care for Hemophilia and Bleeding Disorders:

4:15pm: Patient Perspective: Hear about the personal experiences and concerns navigating healthcare while living with a bleeding disorder.

#### 4:30 - 5:30pm: The Hemophilia Treatment Centers in South Carolina

Meet the South Carolina Hemophilia Treatment Centers and the healthcare teams who treat patients who have a diagnosis of Hemophilia, von Willebrand Disease, platelet disorders, and other rare factor deficiencies.

#### **Presenters:**

Prisma Health Midstate: Stephanie Phillips Ambrose, DO,

Robin Jones, MSN, NHA, CPN, CPHON

Prisma Health Upstate: Leslie Gilbert, MD MSCI

Medical University of South Carolina: Shawn Jenkin Children's Hospital: Shayla Bergmann, MD

5:30pm- Closing and Q & A

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#### **Saturday:**

9:00am: The Hemophilia and Rare Bleeding Disorders Awareness Series

(attendees will receive (1) CEU (55-minute program)

Presented by: Dr. Joanna Davis, MD, University of Miami, Hemophilia Treatment Center

The Hemophilia Awareness Series has been educating medical personnel across the United States for 13 years. The program will educate physicians, nurses, and the medical community on the treatment of Hemophilia in hospitals and emergency rooms.

10:10am: Shemophilia: Women with Bleeding Disorders

<u>Presented by:</u> Dr. Sarah O'Brien, MD, Associate Professor of Pediatrics at Nationwide Children's Hospital This session will provide education on symptomatic carriers and women who have a diagnosis of hemophilia. Learn about the benefits and challenges of diagnosis, how to treat, and how to ensure the patients voice is heard.

11:10am: Emergency and Orthopaedic Issues for Persons with Bleeding Disorders

#### Presented by: Sue Geraghty, RN, MBA

This session is designed to help emergency room personnel and those who practice in the field of orthopaedics understand what the issues are that they may face in dealing with persons with bleeding disorders.

#### **About Accreditation**

In support of improving patient care, this activity has been planned and implemented by Dannemiller and Hemophilia Foundation of Greater Florida. Dannemiller is jointly accredited by the Accreditation Council for Continuing Medical Education (ACCME), the Accreditation Council for Pharmacy Education (ACPE), and the American Nurses Credentialing Center (ANCC), to provide continuing education for the healthcare team.

About the Hemophilia of South Carolina: HSC is a 501(c)3 nonprofit patient advocacy organization representing all South Carolinians with bleeding disorders. Our mission is to raise awareness for and advocate on behalf of persons with bleeding disorders and their families; providing education and supportive services; and promoting ongoing research to improve the quality of life for those affected. Our vision is to be recognized as a leading organization providing valued services; as ambassadors of public outreach to enlighten and foster an understanding of what matters most to those affected by hemophilia and bleeding disorders; and to be our community's first choice in partnership to achieve their highest potential through empowerment, connection to their community and being part of the solutions that affect them the most, until a cure is achieved.

Register Today: tinyurl.com/SCBD2021MS