

REGISTRATION

Name: _____
Address: _____
City: _____
State: _____ ZIP: _____
Phone: _____
Institution: _____

Registration Fee:

Physicians: \$240
Nurses: \$100
General: \$100
Social Workers: \$100
Students: \$50

Last day for registration: March 20, 2025
Make check or money order payable to
Sickle Cell Foundation Conference.

Return to:

The Sickle Cell Foundation
1336 Vickers Road
Tallahassee, Florida, 32303

For electronic registration and payment
by debit or credit card please visit
sicklecellfoundation.org to register and
submit payment.

For Information: Call 850-222-2355



A group of rooms are reserved for guests of the conference at the Holiday Inn. When making reservations, mention the Sickle Cell Foundation to obtain the special conference rate of \$119 per night.

PLACE
STAMP
HERE



**The Sickle Cell Foundation
of Tallahassee, Florida**

Presents

**SICKLE CELL
DISEASE:
*Call to Action,
From Policy to
Implementation*
APRIL 5, 2025**

**Holiday Inn
2003 Apalachee Parkway
Tallahassee, Florida 32301
850-792-8300**

CME, CNE, CEU Application Submitted

Purpose: to provide the most current information on the treatment of and care of patients with Sickle Cell Disease....

COURSE OBJECTIVES

At the end of the conference each participant will be able to :

1. Discuss the rationale for implementation of the Community Education Project.
2. List three best practices for the approach to delivery of care to the Sickle Cell Population.
3. Analyze the need for establishment of policy related to the care of individuals with Sickle Cell disease.
4. State three actions taken by the legislature to improve delivery of care to the sickle cell population.
5. Discuss actions that can be taken on the local, regional, and national level to assist in establishing best practices for delivery of care to the sickle cell population.
6. State three activities that contribute to the development of policy for the approach to the treatment of patients with sickle cell disease.

CONFERENCE SCHEDULE

9:00 am – 9:05 am **Welcome and Introduction**
Alfreda Blackshear, MD

9:05 am – 9:10 am **Greetings**
Velma Stevens, MSW, LSW

9:10 am – 9:15 am **The Occasion**
Latara Lampkin, PhD

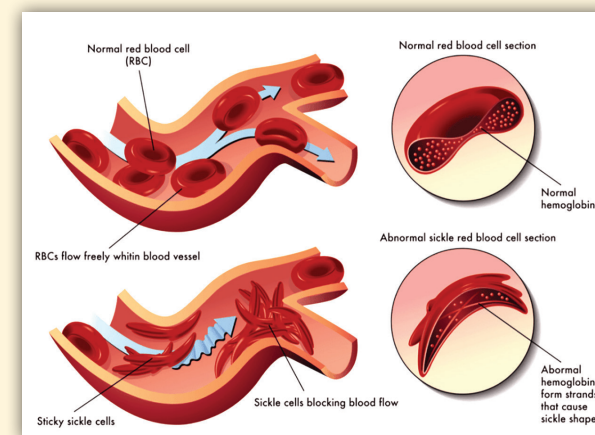
9:15 am – 9:20 am **Instructions**
Mildred D. Fennal, PhD, RN, CNS

9:20 am – 10:20 am **The Trail We Blazed and Why**
Mildred D. Fennal, PhD, RN, CNS

10:20 am – 10:30 am **BREAK**

10:30 am – 11:30 am **Best Practices for Delivery of Care to the Sickle Cell Patient**
James Taylor, MD

11:30 am – 12:30 pm **What We Need is Policy**
*Mapillar Dahn, AD/
Latara Lampkin, PhD*



12:30 am – 1:30 pm **LUNCH**

1:30 pm – 2:30 pm **The Role of the Sickle Cell Disease Association of America in Assisting with Policy Development for Sickle Cell Disease**
Regina Hartfield, BA

2:30 pm – 2:45 pm **BREAK**

2:45 pm – 3:45 pm **Legislative Advances in Sickle Cell Disease Policy (PANEL DISCUSSION)**
Rep. Danny K. Davis, PhD

3:45 pm – 4:45 pm **Strategic Action for Policy Development and Implementation**
Latara Lampkin, PhD

5:00 pm – 5:30 pm **Evaluations and Closing Remarks**