



1

---

---

---

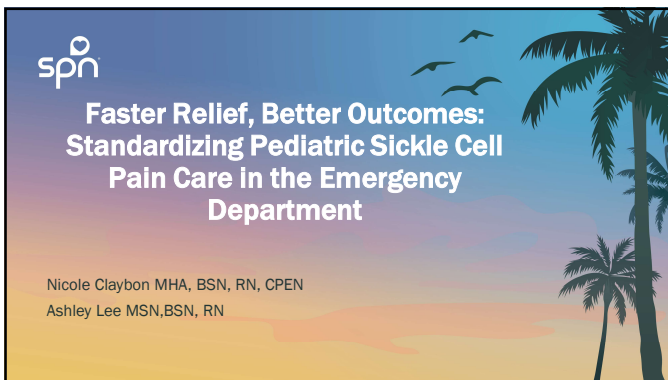
---

---

---

---

---



2

---

---

---

---

---

---

---

---



3

---

---

---

---

---

---

---

---

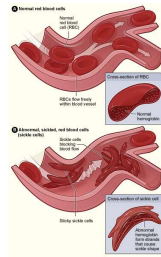
### Pathophysiology of Sickle Cell Pain



Cause of Pain: In sickle cell disease (SCD), red blood cells become rigid and shaped like crescents or "sickles." These cells can clump together and block blood flow in small vessels, leading to Vaso-occlusion. This lack of oxygen causes severe pain in bones, joints, chest, and abdomen.

#### Triggers that Worsen Pain:

- Infection or fever increases sickling.
- Dehydration thickens the blood and worsens clumping.
- Cold exposure or stress causes blood vessels to constrict, raising risk of blockage.
- Hypoxia (low oxygen) from things like high altitude or asthma can also precipitate crises



4

---

---

---

---

---

---

---

---

---

---

### Why Awareness Matters



- ~100,000 people in the U.S. live with sickle cell disease
- Vaso-occlusive crises are the most frequent cause of ED visits in this population
- Most common inherited blood disorder in the U.S.
- National evidence shows children often wait more than 90 minutes for their first pain medication, exceeding the benchmark of 30 minutes



5

---

---

---

---

---

---

---

---

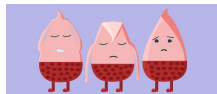
---

---

### Identified Gaps in ED Pain Management



- Promptly initiate interventions following the organization's Sickle Cell protocols
- Improve collaboration within the multidisciplinary team to achieve administering pain meds within 30 minutes
- Identify knowledge gaps and deficiency trends.
- Provide education to staff, utilizing engaging activities, to improve patient care and outcomes.



6

---

---

---

---

---

---

---

---

---

---

### Purpose of this Initiative

- Improve patient experience
- Reduce delays in pain management for pediatric SCD patients
- Decrease admissions
- Improve outcomes for patients through timely identification and intervention
- Increase staff engagement and compliance to standardize sickle cell pain protocols
- Meet the benchmark of initiating appropriate pain management within 30 min of arrival

---

---

---

---

---

---

---

---

7

### Timeliness Goals

Step	Target Time
Door to Room	< 15 minutes
Door to Orders	< 20 minutes
IV Placement	≤ 20 minutes
Door to First Dose	≤ 30 minutes
Pain Reassessment	Every 15-30 min

💡 If no IV w/in 15 min → Give **IN fentanyl** per order

---

---

---

---

---

---

---

---

8

### Sickle Cell Pain Improvement Initiative

- **Direct Rooming:** Assign ESI 2 & room immediately
- **First Pain Med <30 min** (NHLBI, 2014)
- **Verbal Order Set:** Morphine 0.1 mg/kg IV q20min x3 + adjuncts
- **Pain Reassessments:** Every 20 min (wake if asleep)
- **IV Access & Escalation:** Prioritize IV; if delayed, use **IN fentanyl**
- **IV Access Algorithm** – Streamlined process and escalations to minimize delays in venous access.
- **Communication:** Notify provider after 2nd morphine dose; escalate to charge nurse & attending
- **Reminder:** Nurses should also prompt providers to order **Hycet/Norco + Motrin** with morphine

---

---

---

---

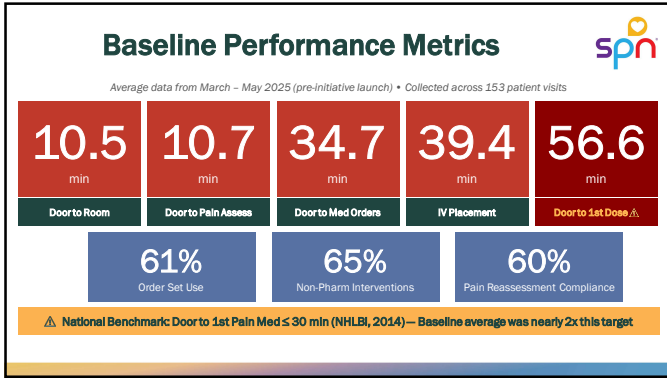
---

---

---

---

9



10

---

---

---

---

---

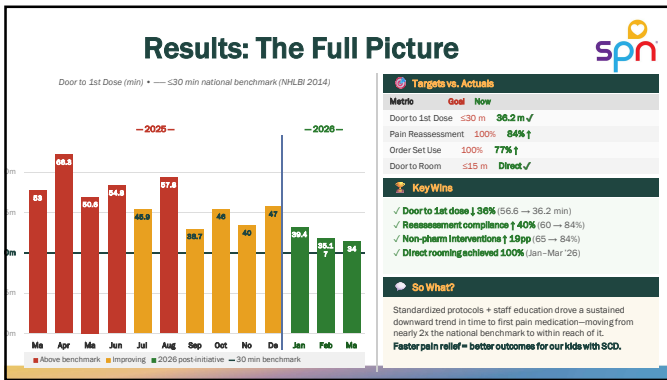
---

---

---

---

---



11

---

---

---

---

---

---

---

---

---

---

### Sustaining Sickle Cell Pain Engagement Through Innovation and Collaboration

#### Strategies that Build Lasting Impact

- Ongoing Evaluation & Coaching – Monthly audits, coaching sessions, and continuous staff education
- Collaborative Practice Refinement – Embedding Sickle Cell Pain strategies into unit-based QI cycles

12

---

---

---

---

---

---

---

---

---

---



