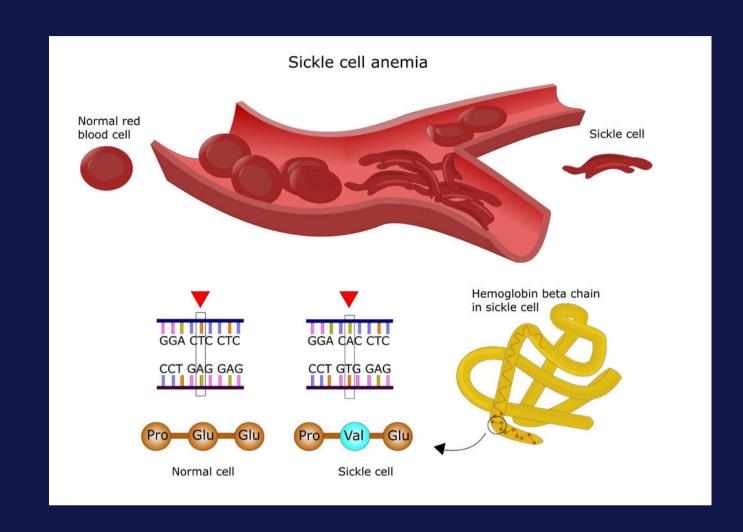


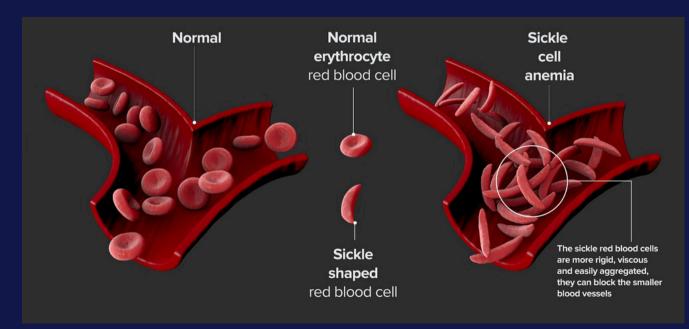
# EVALUATION OF THE NEUROPATHIC PAIN COMPONENT IN SICKLE CELL DISEASE

Maya Mohan

#### INTRODUCTION

- Sickle cell disease: inherited hemoglobin disorder
- Sickle cell mutation: misshapen red blood cells, blockage of blood vessels, pain and organ damage
- Acute and Chronic Pain in sickle cell disease (Ballas 2012)
- Chronic pain: nociceptive and neuropathic
  - Nociceptive: tissue damage
  - Neuropathic: Damage or disfunction of the nervous system
- Neuropathic pain assessment for appropriate pain targeting plan
- Pharmacological and non-pharmacological therapies and biopyschosocial approach
- Biopyschosocial model: framework recognizing biological, pyschological, and social factors (Childerhose 2023)



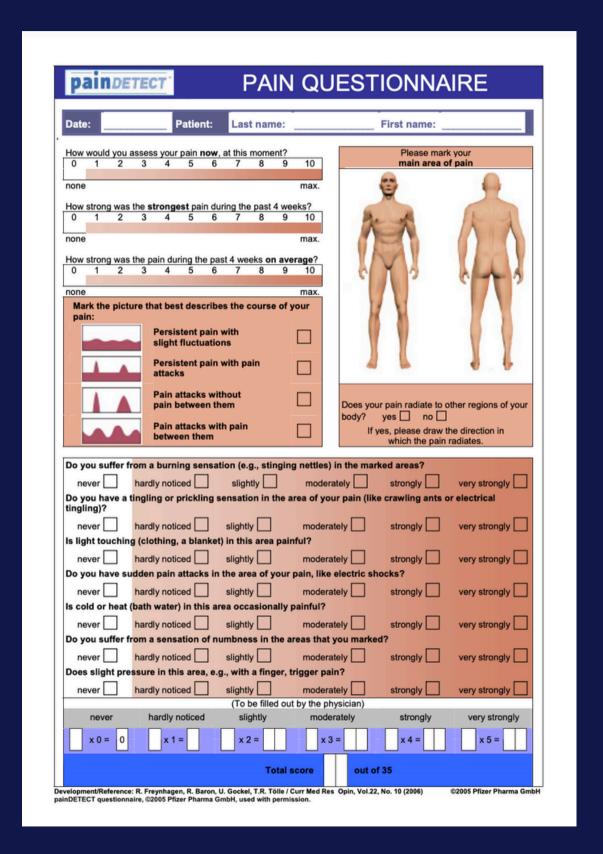


## BACKGROUND/CONTEXT/SIGNIFICANCE

- Descriptive study: neuropathic component in adults with sickle cell disease using painDETECT questionnaire
- Transition from acute to chronic pain and variability of chronic pain not well understood (Cregan 2022)
- Management of chronic pain is therefore inadequate
- Not enough research on the prevelance of neuropathic pain and its effect on the complexixty and severity of pain (Brandow 2014, Glaros 2020)
- Lack of standardized approach to identification, evaluation, and management of neuropathic pain (Brandow 2020)

#### **METHODS**

- painDETECT questionnaire (Freynhagen 2006)
- Adults with sickle cell disease > 18 years
- Neuropathic pain screening tool: 12 questions to differentiate neuropathic vs. non-neuropathic pain
- Total score: 0 to 38
  - ≥ 19 → Definitive Neuropathic Pain
  - 13 to 18 → Probable Neuropathic Pain
  - ≤ 12 → No Neuropathic Pain



#### **RESULTS**

- Analysis of 24 individuals with sickle cell disease
- 66.7% female, 33.3% male
- Median Age: 37 years
- Mean Pain Score
  - At the time of data collection: 5.2
  - Past 4 weeks: 6.8
  - Average strongest pain: 5.3
- 62.5%: Pain Attacks (with or without pain between them)
- 87.5%: Radiation of pain

### FIGURE I: DISTRIBUTION OF PAIN BEHAVIOR PATTERN AMONG STUDY PARTICIPANTS

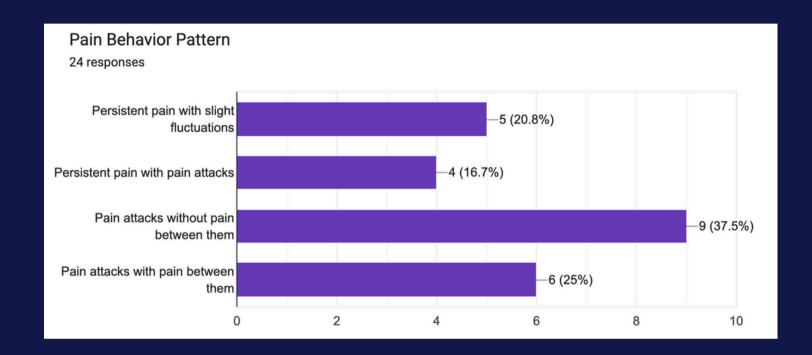


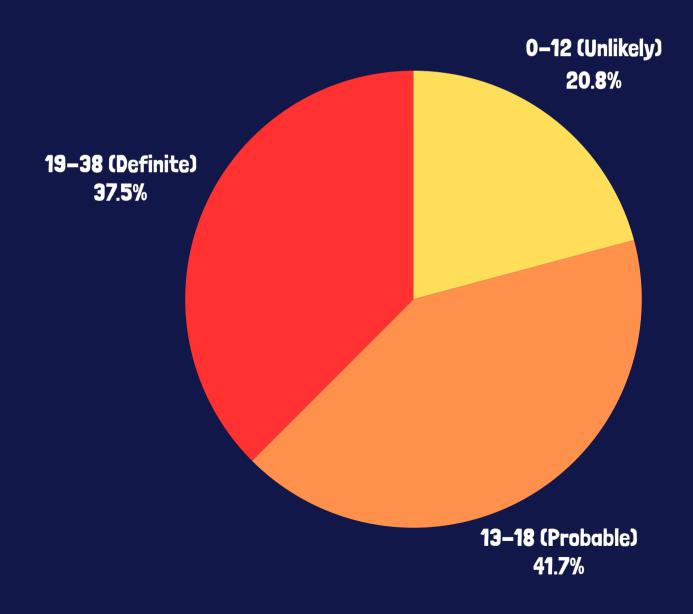
FIGURE II: DISTRIBUTION OF SITES OF PAIN AMONG STUDY
PARTICIPANTS

Site of Pain	Number of Individuals (out of 24)	Percent of Population	20				
Back	17	70.833%	10				
Upper Extremity (Shoulders, Chest, Abdomen, Elbows, Arms)	15	62.5%	5				
Lower Extremity (Hips, Legs, Feet, Knees, Toes)	17	70.833%	0	Back Upper Extremi	Lower Extre	micy, Hips	Head
Head	3	1.25%		John	LOWELL		

#### **CONCLUSIONS AND ANALYSIS**

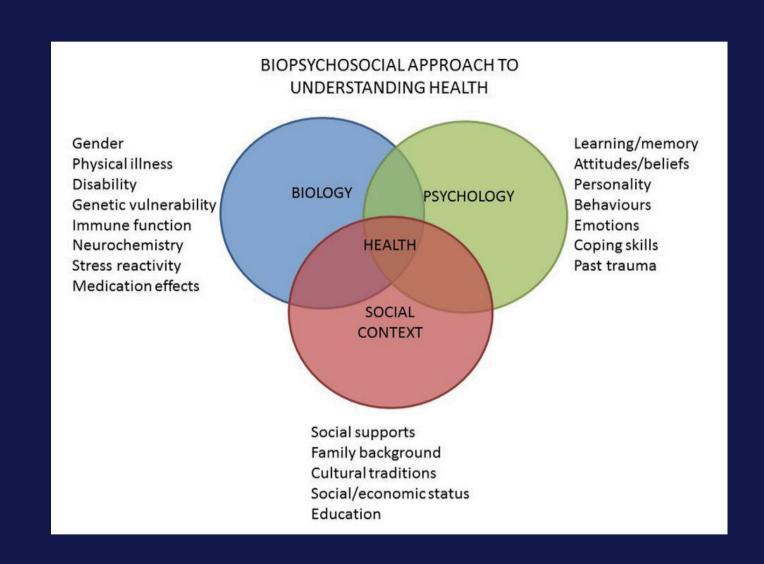
- Primary Outcome Measure
  - 79.2%: Evidence of Neuropathic Component (> 13)
  - 37.5%: Definitive Neuropathic Component (> 19)
  - 41.7%: Probable Component (13 to 18)
- Secondary Outcome Measure
  - Association between total score, age, and gender: no correlation

## FIGURE III: DISTRIBUTION OF SCORES FROM THE PAINDETECT QUESTIONNAIRE AMONG STUDY PARTICIPANTS



#### IMPLICATIONS AND NEXT STEPS

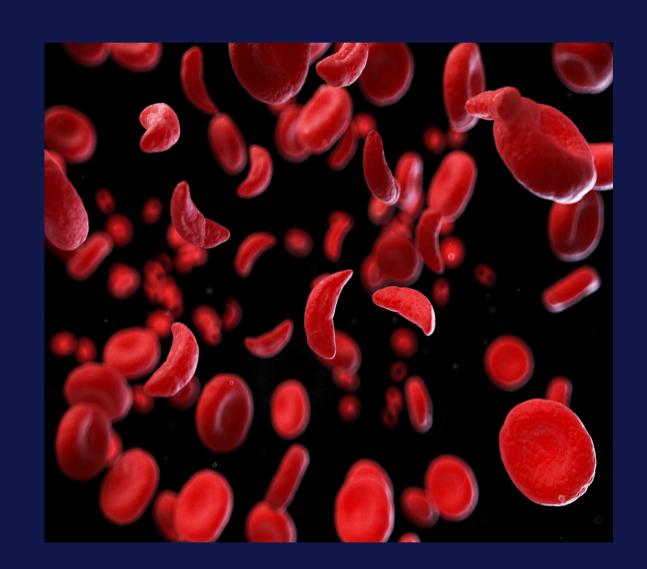
- Identification of the neuropathic component of pain can be used to help plan pain management strategies for individuals. (Brandow 2014, Brandow 2020)
- Recognition of the different component of pain will result in different pain management strategies
- Promotion of a biopsychosocial approach (Childerhose 2023) in addition to pharmacologic and non-pharmacologic therapies.



ASPECTS OF THE BIOPSYCHOSOCIAL MODEL (ONG 2024)

#### CONCLUSION

- The optimal treatment for sickle cell disease: pharmacological, non pharmacological and integrative therapeutic interventions
- Identification of the prevalence of neuropathic pain:
   education, empowerment, and advocacy for individualized
   pain plans with chronic pain component
- Incorporation of a biopsychosocial model: recognition that biologic, neuropsychosocial and socio-environmental elements play a role in pain-related processes
- Enhancement of patient centered care for sickle cell disease



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# THANK YOU