The Evolution of the COVID-19 Pandemic in Pediatric Patients with Sickle Cell Disease: From Alpha to Omicron

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Introduction

- In March 2020, COVID-19 was declared a global pandemic caused by SARS-CoV-2
- Sickle Cell Disease (SCD) was classified as a risk factor for • severe COVID-19 disease in both adult and pediatric patients.
- There was a need for consistent management • guidelines and strategy in treating SCD patients with • COVID 19.
- Complications of SCD treated in the ED include vasoocclusive crises (VOC), or pain crises, and acute chest syndrome (ACS).

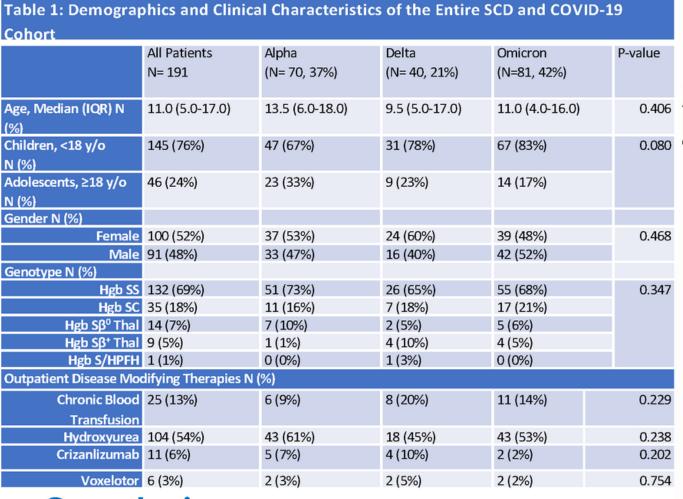
OBJECTIVE:

 To describe the one-year experience of demographics, clinical presentation and management of COVID-19 in children, and young adults with SCD at Children's National Hospital (CNH).

Methods

- Single-center, observational cohort study, with the creation of a SCD COVID-19 registry.
- Children (age <18 years) and young adults (age ≥18 years with SCD and confirmed SARS-CoV-2 positive PCR.
- Evaluated at CNH between March 2020 – January 2023.
- Data collected from the electronic medical record:
- Demographics
- COVID 19 Disease severity (per SECURE registry criteria) and transmission patterns
- Laboratory markers

Results



Conclusions

- In our cohort, we saw no mortality within pediatric and young adult SCD patients w/ COVID-19
- Similar morbidity patterns (ACS, Pain/VOC) between SCD pediatric and young adult patients w/COVID-19.
- Hospitalized patients were more likely to have fever, oxygen requirement, VOC and ACS.
- ACS patients were more coagulopathic, anemic, and hypoxic requiring blood transfusions.
- No Hydroxyurea (HU) use was associated with being hospitalized and higher acuity
- Future studies will characterize long COVID symptoms in pediatric and young adults w/ SCD.

Results

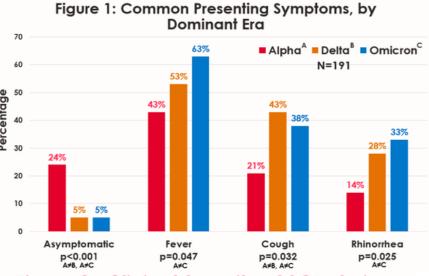
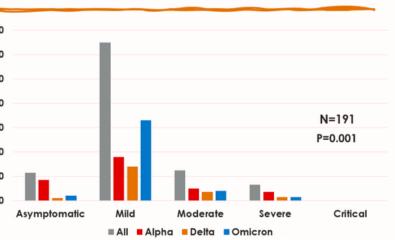


Figure 2: Clinical Severity of SCD during COVID-19 Period



*Adapted from the SECURE SCD rea





Spousors

