Sickle Cell Disease (SCD) and COVID-19 – A Case Series

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Case Report

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Abstract

The ongoing pandemic of COVID-19 that started in the Hubei province of China in late December 2019, caused by severe acute respiratory syndrome corona virus-2 (SARS CoV-2). Globally millions affected by the disease so far. The risk of COVID-19 severity and its complications increases with age and other comorbidities. The course of SARS-CoV-2 infection or its related complications has yet to be established in patients with sickle cell disease (SCD), once more evidence is available. It is clear from the available data that the course of COVID-19 in patients with SCD is mild to moderate, seldom severe, and rarely fatal. Herein we report three known cases of SCD with confirmed COVID-19, in whom the course of the disease was mild to moderate and uncomplicated with uneventful recoveries.

Introduction

The current pandemic of COVID-19 caused by SARS-CoV-2 affects millions worldwide. As of November 26, 2020, worldwide confirmed infections are over 60 million in 191 countries/territories¹. The disease brought challenges to almost every sector of life, specifically health. The more vulnerable individuals are aged and with comorbidities². The most challenging aspect of healthcare is the care of patients with comorbidities that necessitate balancing the care needed with their exposure to SARS CoV-2 infection.

Sickle cell disease (SCD) an autosomal recessive disorder due to a single point mutation of glutamic acid by valine at position six. Patients with SCD pose a management challenge for healthcare due to the complexity of the disease and the scarce published data on pathophysiological interaction of SARS CoV-2 infection in sickle cell disease. Herein, we report a case series of three known patients of SCD with confirmed infection of SARS CoV-2.

Case Presentations

Case 1:

A 23year old female, a known patient of SCD with HbSS, presented to the emergency department with typical findings of vaso-occlusive crisis (VOC) and nasal congestion. There was no history of cough or fever. The patient's history is suggestive of SCD with repeated VOC. Due to the current COVID-19 pandemic, clinical evaluation was done under standard precautions and was unremarkable. Vitals were; pulse, 86/min, BP, 117/78mmHg, and respiratory rate, 20/min. Laboratory workup shows; Hb 10.3g/dL, WBCs $9.1 \times 10^3/\mu\text{L}$, $375 \times 10^3/\mu\text{L}$. Oxygen saturation was 96% with 3lit supplemental oxygen by face mask. IV fluids initiated in ED with IV morphine 0.1mg/kg. After two hours of management in the ED, the patient shifted to an isolation unit for close observation and management. Biochemistry result shows s. urea, 2.34mmol/L, creatinine, 60mmol/L, Na^+ 135mmol/L and K^+ , 3.2mmol/L. Nasopharyngeal swab, taken for RT-PCR, which came out positive the next day. Medications includes hydroxyurea as the patient's medication, IV ceftriaxone, paracetamol, Vitamin-D, and calcium carbonate. The pain severity

improved the next day, and the patient was discharged home after two days with specific COVID-19 recommendations.

Case 2:

A 30year old male, a known patient of SCD with HbSS, presented to the emergency department with fever and generalized body aches since last night but no history of cough. Under the standard protocol, clinical evaluation was unremarkable with vitals as; Temp 39C, pulse 91/min, BP 115/75 mmHg. Oxygen saturation was 95% with 3lit oxygen by face mask. Initial laboratory workup in the ED shows; Hb 13g/dL, WBCs $8.7 \times 10^3/\mu\text{L}$, $305 \times 10^3/\mu\text{L}$. With early initiation of IV fluids and IV analgesics (morphine) with oxygen supplementation, the patient's symptoms improved. After initial management in the ED, the patient shifted to the isolation ward with standard protocol. COVID-19 test requested, a nasopharyngeal swab taken by the assigned team for real-time polymerase chain reaction (RT-PCR) with a positive report a day after. Biochemistry workup was within normal limits. CXR showed no positive findings. The patient was put on regular doses of IV ceftriaxone, IV morphine, Tab favipiravir, calcium carbonate, and Vitamin-D as per protocol. The patient's symptoms improved with no further intervention and discharged home on 3rd day of admission with standard recommendations.

Case 3:

A 21year old male patient, a known case of SCD with HbSS with a history of contact to COVID-19 positive patient, presented to the primary care center with a history of sore throat and two episodes of loose motion. There was no history of cough or fever. Due to contact with a COVID-19 patient, RT-PCR requested. The patient was advised for quarantine after primary care management, to be followed, if needed. The test result came positive, a day after. The patient followed for the next ten days with no mishap.

Discussion

The present pandemic of COVID-19 caused by severe acute respiratory syndrome corona virus-2 (SARS CoV-2) is a public health crisis of the century. The disease affected millions globally. Individuals with comorbid conditions are more vulnerable than healthy ones. The course of COVID-19 varies from asymptomatic to more severe. Despite the comorbid condition of SCD, the patient clinical course is mild, and the severity is less in this group.

Nur et al³. reported two SCD patients with HSS. Both were positive for COVID-19, and their presenting symptom was pain. Both patients recovered well. Three patients of one family reported by Abdulqader et.al⁴, the symptoms ranged from asymptomatic to VOC and ACS in one patient with the un-eventful recovery of all the three.

Mass screening in Bahrain⁵ reports a total of 378 patients with SCD, 1% of the screened population. A total of six patients were positive for COVID-19. All six patients were HSS genotype, of whom three

remained asymptomatic, two patients had mild symptoms, and only one required supplemental oxygen. All patients recovered well. Menapace and Thein⁶ concluded the same observation of less severe outcome of COVID-19 in SCD patients, and the usual presentation was pain not fever.

There is conflicting evidence of COVID-19 severity in SCD. In the UK⁷, a massive survey over 4-weeks, there were 195 confirmed or clinically suspected cases of COVID-19, the majority of the cases (n=166) were patients with SCD. Severe course of disease and mortality recorded in aged patients and not in children.

Pre-exposure prophylaxis is a hallmark of patient management with SCD during the pandemic of COVID-19. Patients are advised and encouraged to adhere strictly to social distancing, hand hygiene, and avoiding unnecessary exposure to public places, and also to adhere to their disease-modifying medications to minimize the frequencies of VOC, thus subsequently hospital visits.

From the limited published data, it is not clear whether SCD increases the possibility of SARS CoV-2 infection but, it is clear from the available data of case reports that the course of COVID-19 was mild-moderate and rarely severe. The protective factors in SCD patients for SARS CoV-2 infection need to be explored with a clearer understanding of the relationship between SCD and COVID-19 once more evidence is available.

Conclusion

The course of COVID-19 in these patients was mild to moderate with un-eventful recovery. Considering SCD patients with concurrent COVID-19 as high-risk individuals and should be managed with established protocols.

Declarations

Ethical approval

Not applicable

Consent:

Written informed consent was taken from patients for publication of their cases.

Conflict of Interest:

The authors declare that they have no conflict of interest.

References

1. The Johns Hopkins University of Medicine. COVID-19 dashboard by the Center for Systems Science and Engineering at Johns Hopkins University. <https://coronavirus.jhu.edu/map.html> Date accessed: November 26, 2020.
2. Hussain, F.A., Njoku, F.U., Saraf, S.L., Molokie, R.E., Gordeuk, V.R. and Han, J. COVID-19 infection in patients with sickle cell disease. *Br J Haematol*, 2020;189: 851-852. <https://doi.org/10.1111/bjh.16734>
3. Nur, E, Gaartman, AE, van Tuijn, CFJ, Tang, MW, Biemond, BJ. Vaso-occlusive crisis and acute chest syndrome in sickle cell disease due to 2019 novel coronavirus disease (COVID-19). *Am J Hematol*. 2020; 95: 725– 726. <https://doi.org/10.1002/ajh.25821>
4. Abdulqader Al-Hebshi, Mohammed Zolaly, Amer Alshengeti, Ghaya Al Qurainees, Sofyan Yamni, Naif Hamadan, Turki Alwasaidi. *Pediatric Blood & Cancer*. 2020;67:e28547. DOI: 10.1002/pbc.28547 <https://doi.org/10.1002/pbc.28547>
5. Abdulkarim AbdulRahman, Salman AlAli, Omar Yaghi, Mohammed Shabaan, Sameer Otoom, Stephen L. Atkin, Manaf AlQahtani. COVID-19 and Sickle Cell Disease in Bahrain, *International Journal of Infectious Diseases* (2020), <https://doi.org/10.1016/j.ijid.2020.09.1433>. <http://www.sciencedirect.com/science/article/pii/S1201971220321494>
6. Laurel A. Menapace and Swee Lay Thein. COVID-19 and sickle cell disease. *Haematologica*.2020; 105(11): 255398 doi: 10.3324/haematol.2020.255398
7. Telfer P, de la Fuente J, Sohal M, Brown R, Eleftheriou P, Roy N, Piel FB, Chakravorty S, Gardner K, Velangi M, Drasar E, Shah F, Porter JB, Trompeter S, Atoyebi W, Szydlo R, Anie KA, Ryan K, Sharif J, Wright J, Astwood E, Nicolle CS, Webster A, Roberts DJ, Lugthart S, Kaya B, Awogbade M, Rees DC, Hollingsworth R, Inusa B, Howard J, Layton DM. Real-time national survey of COVID-19 in hemoglobinopathy and rare inherited anemia patients. *Haematologica* 2020;105(11):2651-2654; <https://doi.org/10.3324/haematol.2020.259440>.