CASE REPORT

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COVID-19 in a pregnant patient with beta-thalassemia major: A case report

Yousef Mohammed Ali Hailan¹ Gamal Saved^{2,3} Mohamed A. Yassin⁴

Abstract

¹Internal Medicine Department, Hamad Medical Corporation, Doha, Qatar

²Women Wellness and Research Center Medical, Obstetrics/Gynecology, Hamad Medical Corporation (HMC), Doha, Qatar

³Clinical Department, College of Medicine, QU Health, Qatar University, Doha, Qatar

⁴National Center for Cancer Care and Research, Hamad Medical Corporation, Doha, Qatar

Correspondence

Yousef Mohammed Hailan, Internal Medicine Department, Hamad General Hospital, Hamad Medical Corporation, Al Rayyan Street, Doha, 00974, Qatar. Email: yhailan@hamad.qa; yousefhailan@ live.com

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1 **INTRODUCTION**

Beta-thalassemia major, a prevalent disease, is caused by severely reduced or absent beta-globin production. Chances of pregnancy have increased significantly since the introduction of hypertransfusion and iron chelation therapies. We report a case of a 35-year-old Lebanese pregnant lady with a background of beta-thalassemia major who was diagnosed with COVID-19 infection (cycle threshold value 18) during her 23rd gestational week. Unfortunately, the pregnancy outcome was unfavorable as it was complicated by intrauterine fetal death. To our knowledge, this is the first report of such a case.

The coronavirus disease (COVID-19) is caused by the novel severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2). COVID-19 has a wide range of presentations, and its severity varies from asymptomatic disease to

crease their chances of normal pregnancy and fewer complications and more favorable outcomes. **KEYWORDS**

Further studies are needed on this unique population to better manage them and in-

beta-thalassemia major, COVID-19, pregnancy, stillbirth, thalassemia, transfusion-dependent

life-threatening sepsis.¹ Since it surfaced in Wuhan, China, in December 2019 and was announced as a pandemic by the World Health Organization (WHO) in March 2020, it has resulted in over 126 million confirmed cases and more than 2.7 million deaths globally unto 30 March 2021.^{2,3} Previous studies revealed that droplets, contact, aerosol, and fecal-oral transmissions are the main transmission routes in COVID-19 infection.⁴ Vertical transmission is believed to be less of a concern.⁵ Although many publications have discussed the association between many comorbidities and the severity of COVID-19 infection, data on the COVID-19 and hemoglobinopathies are still limited.6-8

Variants of thalassemia produce a wide range of clinical manifestations. Homozygotes for β-thalassemia may develop either thalassemia major or thalassemia intermedia. β-thalassemia is caused by partial or total reduction in the

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 β -globin chains in the HbA molecule. Among Arab populations, the carrier rates range from 1% to 11%, and the most frequent mutation is IVS-1-110 (G > A).⁹ Furthermore, Khan et al have identified six unique β -thal mutations in six Arab countries.¹⁰

Beta-thalassemia major manifests in infancy with a constellation of symptoms including pallor, jaundice, and failure to thrive; physical examination findings of hepatosplenomegaly, frontal bossing, and thalassemic facies; and laboratory investigations consistent with a microcytic anemia with hemoglobin <7 g/dL; and hemolysis.¹¹

The primary treatment of this type of anemia is with a regular transfusion schedule targeting a pretransfusion hemoglobin level between 9 and 10 g/dL, preferably transfusions of washed, leukocyte-depleted red blood cells to reduce the incidence of reactions, along with addressing the complications as appropriate, namely endocrinopathies such as hyperadrenalism and abnormalities in glycemic control and insulin-like growth factor-1(IGF-1).¹¹⁻¹⁶

Unlike patients with alpha-thalassemia, pregnancy in women with beta-thalassemia major was associated with unfavorable outcomes until after the introduction of hypertransfusion and iron chelation therapies in the late 1970s.¹⁷

We describe a case of a woman with a beta-thalassemia major who acquired a COVID-19 infection during her pregnancy and the outcome of the pregnancy.

2 | CASE REPORT/CASE PRESENTATION

A 35-year-old Lebanese female patient gravida 4 para 1 presented to the hospital with fever and dry cough for 3 days. She is known to have beta-thalassemia major on regular transfusions every 3 weeks, and the last transfusion was 5 days before this presentation. She also has a history of cholecystectomy and splenectomy. She is not known to have any allergies. She was taking aspirin and deferasirox at home. She is a teacher, and both of her parents are carriers of betathalassemia trait. Otherwise, family and social history are noncontributory.

Physical examination was nonsuggestive, and admission laboratory investigations (shown in Table 1) showed mild leukocytosis, hemoglobin (Hb) at target, normal renal function, slightly elevated liver enzymes, and markedly elevated ferritin. Chest XR was reported normal.

The evaluation revealed that she has a mild COVID-19 infection with a cycle threshold value of 18. She was pregnant in week 27, as calculated from the last menstrual period (27 September 2020). It was confirmed later by ultrasound (US) to be a single viable fetus aged 23 weeks and 2 days. Upon admission, she was seen by multiple specialties, primarily infectious disease, internal medicine, hematology, and

ABLE 1 Admission laboratory invest	igations
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Detail	Value w/Units	Normal range	
WBC	$13.84 \times 10^{3/uL}$	4.00-10.00	
RBC	$4.1 \times 10^{6/uL}$	3.8-4.8	
Hgb	11.6 gm/dL	12.0-15.0	
Hct	34.4%	36.0-46.0	
MCV	84.4 fL	83.0-101.0	
MCH	28.3 pg	27.0-32.0	
MCHC	33.6 gm/dL	31.5-34.5	
RDW-CV	14.3%	11.6-14.5	
Platelet	$322 \times 10^{3/uL}$	150-400	
MPV	10.5 fL	7.4-10.4	
Absolute Neutrophil Count Auto# (ANC)	$12.4 \times 10^{3/uL}$	2.0-7.0	
Lymphocyte Auto #	$0.6 \times 10^{3/uL}$	1.0-3.0	
Monocyte Auto #	$0.6 \times 10^{3/uL}$	0.2-1.0	
Eosinophil Auto #	$0.1 \times 10^{3/uL}$	0.0-0.5	
Basophil Auto #	$0.09 \times 10^{3/uL}$	0.02-0.10	
Neutrophil Auto %	89.4%		
Lymphocyte Auto %	4.5%		
Monocyte Auto %	4.7%		
Eosinophil Auto %	0.4%		
Basophil Auto %	0.6%		
Prothrombin Time	10.1 s	9.7-11.8	
INR	1.0		
D-Dimer	>4.40 mg/L FEU	0.00-0.44	
Fibrinogen	4.78 gm/L	1.70-4.20	
APTT	39.6 s	24.6-31.2	
Urea	2.60 mmol/L	2.50-7.80	
Creatinine	27 umol/L	53-97	
Sodium	141 mmol/L	133-146	
Potassium	4.1 mmol/L	3.5-5.3	
Chloride	98.7 mmol/L	95.0-108.0	
Bicarbonate	28.4 mmol/L	22.0-29.0	
Bilirubin T	20.6 umol/L	0.0-21.0	
Total Protein	72 gm/L	60-80	
Albumin Lvl	40.2 gm/L	35.0-50.0	
Alk Phos	128.0 U/L	35.0-104.0	
ALT	52.0 U/L	0.0-30.0	
AST	55 U/L	0-31	
Glu Fasting	4.3 mmol/L	3.3-5.5	
NT pro-BNP	52.8 pg/mL	0.0-130.0	
Troponin-T HS	4.1 ng/L	0.0-14.0	
LDH	188 U/L	135-214	
СК	23 U/L	2-160	

(Continues)

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TABLE 1 (Continued)

Detail	Value w/Units	Normal range
G6PD Screen	Normal	
CRP	35 mg/L	0-5
Procalcitonin	0.20 ng/mL	
Ferritin	2,942 mcg/L	8-252
COVID-19 PCR	Positive	
COVID-19 Average CT	18.08	

obstetrics. As per the local Communicable Disease Center (CDC) COVID-19 management protocol, she is for symptomatic treatment.

On day 4 of admission, she reported reduced fetal movement and the urgent obstetric US reported fetal death. The next day, she underwent misoprostol induction protocol for intrauterine fetal death which was uncomplicated. On day 7, she was discharged from the hospital as COVID-19 PCR became negative and her symptoms settled.

3 | **DISCUSSION/CONCLUSION**

De Sanctis et al published a thorough article in 2019 addressing marital status and paternity in patients with transfusiondependent thalassemia (TDT) and non-transfusion-dependent thalassemia (NTDT).¹⁸ The notable observations in patients with TDT include the following: The majority of the patients have natural conception (78.5%), the most common cause of infertility is dysspermia (13.3%), and the average level of serum ferritin in the year of paternity is 2211.8 ± 181.8 ng/ mL.

The introduction of hypertransfusion and iron chelation therapy has increased the chances for these women for pregnancy and better pregnancy outcomes. The likely mechanism by which pregnancy was highly unlikely in this population is primarily due to anovulation secondary to hypogonadotropic hypogonadism due to iron overload in the hypothalamus and pituitary gland.^{17,19,20} The most recent American College of Obstetricians and Gynecologists recommendations advise pregnancy in women with TDT only to those with normal cardiac function, prolonged hypertransfusion therapy to maintain Hb levels at 10 g/dL, and iron chelation therapy with desferrioxamine.

Iron chelating agents aim to excrete the accumulating iron through feces and/or urine. The currently approved chelators are desferrioxamine (DFO), deferasirox (DFX), and deferiprone (DFP).^{21,22} However, the safety profile for these agents is not well studied in pregnancy, and the usual recommendation is to hold them during pregnancy. Since holding chelating therapy for the duration of pregnancy may have important consequences on women, some researchers prefer to use DFO in the second and third trimesters as it is a large molecule and less likely to cross the placenta.

A recent systematic review on pregnancy and COVID-19 that included a total of 8 studies involving 95 pregnant women and 51 neonates addressing the maternal, obstetric, and neonatal outcomes concluded that contrary to severe acute respiratory syndrome coronavirus (SARS-CoV) and Middle East respiratory syndrome coronavirus (MERS), SARS-CoV-2 does not appear to increase the risk of pregnancy complications.²³ Another publication suggested that a high rate of maternal and fetal complications is seen in infected individuals.²⁴ The most common pregnancy complications in women with COVID-19 were fetal distress, premature rupture of membranes, preterm labor, and post-partum fever.^{5,23}

No data in the literature address the topic of pregnancy in patients with β -thalassemia major in particular or β thalassemia in general in COVID-19 patients. Some of the publications discussing pregnancy in a patient with COVID-19 infection mentioned thalassemia, thalassemia trait, and thalassemia minor in the list of comorbidities in the description of their included patients characteristics.^{5,25-27} However, no details were provided as to the outcomes and course of the pregnancy in this subset of patients.

Our patient is known to have transfusion-dependent thalassemia and was infected with COVID-19. She was managed from a COVID-19 infection point of view as per version 12 of the local CDC recommendations. The recommendation for pregnant females who have positive COVID-19 PCR with uncomplicated upper respiratory tract infection is isolation, either at home or in an isolation facility, and supportive treatment as needed. From the hematology aspect, when she became pregnant, her transfusion schedule changed to receive packed red blood cell transfusions every 2 weeks instead of every 3 weeks.

Despite being managed by a multidisciplinary team, the outcome of the pregnancy was unfavorable. It can be attributed to COVID-19 infection, β -thalassemia major, and iron excess. The placental sample sent for pathological analysis showed early ischemic changes and other features in favor of mild acute chorioamnionitis. Thrombosis is a major complication of COVID-19 infection, and the placenta is not immune.²⁸ Whether the early ischemic changes in the report are linked to COVID-19 infection is uncertain and is debatable.

To our knowledge, this is the first case report that highlights COVID-19 infection in a pregnant patient with betathalassemia major.

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CONFLICT OF INTEREST

The authors have no conflicts of interest to declare.

AUTHOR CONTRIBUTIONS

Yousef Mohammed Ali Hailan and Mohamed A Yassin: performed writing, editing, and final approval of the concept. Gamal Sayed: performed editing and approval of the final version.

ETHICS STATEMENT

The case was approved by Hamad Medical Corporation Research Center with reference number MRC-04-21-352. Written informed consent was obtained from the patient for publication of this case report and any accompanying images.

DATA AVAILABILITY STATEMENT

Data are available on reasonable request.

ORCID

Yousef Mohammed Ali Hailan https://orcid. org/0000-0002-0760-9906

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