

## CLINICAL IMAGE

# Clinical image report of acute Cholangitis in Beta-thalassemia major

Wail Rozi<sup>1</sup>  | Ibrahim Khamees<sup>1</sup>  | Mohamed A. Yassin<sup>2</sup> 

<sup>1</sup>Internal Medicine, Hamad Medical Corporation, Doha, Qatar

<sup>2</sup>Hematology and Oncology, Hamad General Hospital, Doha, Qatar

## Correspondence

Wail Rozi, Internal Medicine, Hamad Medical Corporation, Doha, Qatar.  
Email: wail.rozi@gmail.com

## Funding information

Qatar National Library funded the open access publication fees of this case

## Abstract

Beta-thalassemia is congenital red blood cell disorder. Gallstones is a recognized complication due to recurrent hemolysis. Acute cholangitis is a rare complication might occur in patient with beta-thalassemia. We report a case of acute cholangitis in patient with beta-thalassemia with Gilbert syndrome. We present a case of a young female of Arabic descent with acute abdomen. Workup revealed acute cholangitis with gallstones in the common bile duct. The ERCP was used to extract the stones than cholecystectomy.

## KEYWORDS

acute cholangitis, beta-thalassemia, common bile duct, gall bladder

## 1 | DESCRIPTION

A 37-year-old female patient with a background of beta-thalassemia major presented to the emergency department complaining of right upper quadrant pain associated with nausea and vomiting. Physical examination was consistent with positive Murphy's sign. Blood workup showed a picture of cholestatic injury (Table 1). Ultrasound abdomen (Figure 1) features were suggestive of acute cholecystitis along with dilated common bile having multiple stones

and intrahepatic biliary dilatation. Further imaging with abdomen MRI and MRCP confirmed the previous finding and revealed secondary hemochromatosis (Figure 2).

TABLE 1 Liver function test

Detail	Value w/Units	Normal Range
Bilirubin T	164 umol/L	0–21
Bilirubin D	122 umol/L	0–5
Total protein	62 gm/L	60–80
Albumin level	39 gm/L	35–50
Alkaline phosphatase	403 U/L	35–104
ALT	687 U/L	0–33
AST	682 U/L	0–32

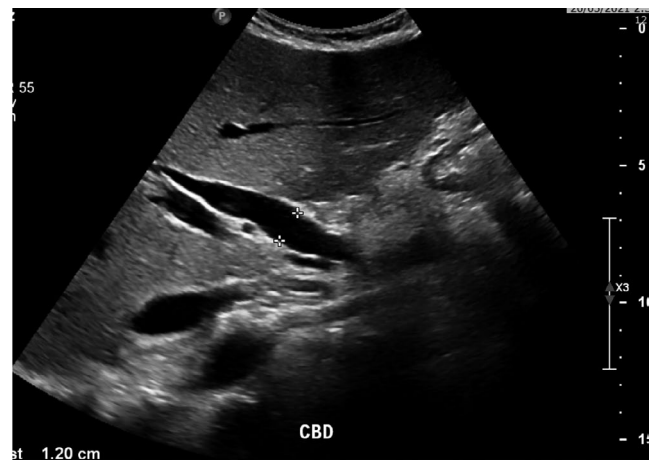
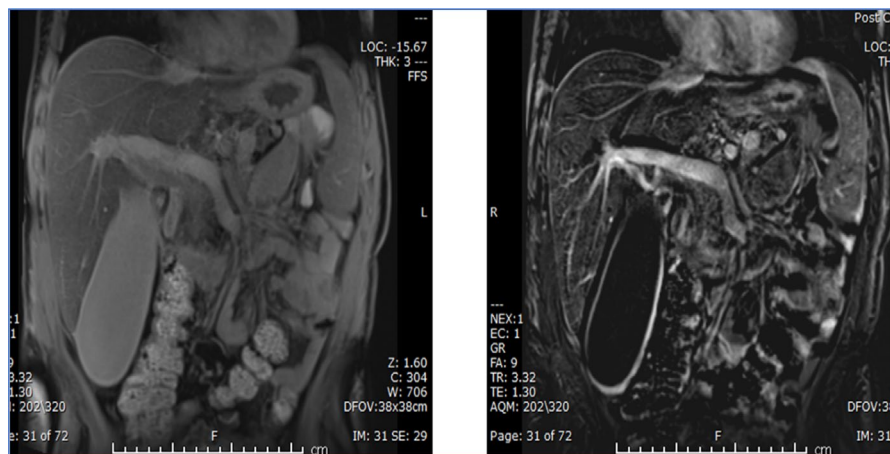


FIGURE 1 Common bile duct is dilated, measuring up to 12 mm. Small stones are seen in the mid-to-distal CBD, the largest measuring 5.5 mm

This is an open access article under the terms of the Creative Commons Attribution-NonCommercial-NoDerivs License, which permits use and distribution in any medium, provided the original work is properly cited, the use is non-commercial and no modifications or adaptations are made.

© 2022 The Authors. *Clinical Case Reports* published by John Wiley & Sons Ltd.



**FIGURE 2** Pre- and post-contrast of T1 sequence, post-contrast enhancement of the ductal system suspicious of cholangitis

Beta-thalassemia is caused by a defect in the synthesis of the beta-globin chain of hemoglobin. Homozygotes for beta-thalassemia can have either beta-thalassemia major or intermedia, which can be distinguished by genetic studies.<sup>1</sup> A frequently encountered complication in beta-thalassemia major is the development of bilirubin gallstones, especially in patients with co-existing Gilbert's syndrome,<sup>1–3</sup> which are usually asymptomatic and do not need any medical intervention. However, the development of cholangitis and cholecystitis is uncommon, and surgical intervention may be indicated. Our patient was treated with ERCP first. Then, she underwent cholecystectomy.

#### ACKNOWLEDGEMENTS

We wish to thank Qatar National Library and the Internal Medicine Residency Program of Hamad Medical Corporation for scientific support.

#### CONFLICT OF INTERESTS

The authors have no conflict of interest relevant to this case.

#### AUTHOR CONTRIBUTIONS

Both authors (Wail Rozi and Mohamed Yassin) contributed equally to the writing and preparation of this article, critically reviewed the initial and the final draft of the manuscript, and approved it for submission. Wail Rozi wrote the initial draft of the manuscript and performed the literature review. The draft was revised and updated by Wail Rozi with support from Dr. Mohamed Yassin. Mohamed A. Yassin supervised, revised, and updated the draft.

#### ETHICAL APPROVAL

Ethics approval and consent to participate. Medical Research Committee at Hamad Medical Corporation approved the case study for publication (MRC-04–21–324).

#### CONSENT


Written informed consent was obtained from the patient for the publication of this clinical image report.

#### DATA AVAILABILITY STATEMENT

Data and materials regarding the case report are available to the editor in chief and can be requested from the corresponding author.

#### ORCID

Wail Rozi  <https://orcid.org/0000-0003-0566-4888>

Ibrahim Khamees  <https://orcid.org/0000-0003-2887-9726>

Mohamed A. Yassin  <https://orcid.org/0000-0002-1144-8076>

#### REFERENCES

1. Khan AM, Al-Sulaiti AM, Younes S, Yassin M, Zayed H. The spectrum of beta-thalassemia mutations in the 22 Arab countries: a systematic review. *Expert Review of Hematology*. 2021;14(1):109-122.
2. Origa R, Galanello R, Perseu L, et al. Cholelithiasis in thalassemia major. *Eur J Haematol*. 2009;82(1):22-25.
3. Soliman A, Yassin M, Al Yafei F, et al. Longitudinal study on liver functions in patients with thalassemia major before and after deferasirox (DFX) therapy. *Mediterr J Hematol Infect Dis*. 2014;6(1):e2014025.

**How to cite this article:** Rozi W, Khamees I, Yassin MA. Clinical image report of acute Cholangitis in Beta-thalassemia major. *Clin Case Rep*. 2022;10:e05705. doi:[10.1002/ccr3.5705](https://doi.org/10.1002/ccr3.5705)