Case Report of the Coincidence of Dubin and Johnson Syndrome and Colon Cancer

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Abstract

Introduction: Dubin and Johnson syndrome (DJS) is autosomal recessive, inherited disorder with no progression to end-stage Liver disease. It results mainly unconjugated hyperbilirubinaemia. Clinical onset of DJS often begins during the teenage years or in early adulthood. There are case reports describe the coincidence of DJS and colon cancer. Brief case report: We describe 27 years old male diagnosed with adenocarcinoma of the colon few years after the clinical onset of DJS. Conclusion: Attention should be taken when evaluating a young adult with DJS and be aware of this risk and consequences of a late diagnosis of colorectal cancer. Keywords: Dubin-Johnson syndrome, Hyperbilirubinemia Colon Cancer.

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INTRODUCTION

In 1954, Dubin and Johnson1 reported 12 cases of chronic jaundice characterized by elevation of both direct and indirect serum bilirubin, autosomal recessive, inherited disorder with no progression to end-stage Liver disease. Both the conjugated and unconjugated form of bilirubin can be elevated in DJS subjects, with the former ranging from 17% to 88% of the total bilirubin with a mean value of 60% [1, 2], in addition to the fluctuating jaundice. DJS subjects may suffer from nonspecific symptoms such as weakness and abdominal discomfort. Urinary coproporphyrin output is normal [11]. Thus, the biliary excretion of anionic dyes including bromosulphophthalein, indocyanine green and cholestintigraphy radiotracers is delayed. Liver histology in DJS shows an accumulation of distinctive melanin-like lysosomal pigment in an otherwise normal liver, which gives the organ a characteristic dark pink or even black colour [1-3]. Clinical onset of DJS is often seen during the teenage years or in early adulthood, which is typical to our patient. In this report, we describe 27 years old male diagnosed with adenocarcinoma of the colon few years after the clinical onset of DJS. However, the relationship between DJS and colon cancer not yet fully understood.

CASE REPORT

This 29 years old male, first time seen in the gastroenterology clinic in June 2013 when he presented by intermittent right hypochondrial pain and jaundice for almost three months. There was no itching, or weight loss. For instance, he has no evidence of gastrointestinal bleeding or Porto systemic encephalopathy. He did not consume alcohol or smoke. He has no family history of jaundice. At this stage his liver profile show: (GTP 91- Bili D 73.1 - Bili T 101.5 - Alk Phos 95- TP 76- Alb 46 - ALT 41- AST 23).

Screen for hepatic autoimmune and viral serology requested and result came back as following: (Hepatitis C Antibody, HBsAg, HBE AG, CMV IgM, ANA-H) ALL negative, AntiHbs >1000.00. Transferrin 2.57 Ceruloplasmin (0, 23). Liver/Kidney Mic antibody (1.25), anti-smooth muscle antibody less than (1:20).

The work up point toward isolated unconjugated hyperbilirubinemia. Therefore, MRCP requested on June 2013 Findings were highly suggestive of early primary sclerosing cholangitis. Other possibility like eosinophilic cholangitis and HIV related cholangiopathy to be consider. Correlation with ERCP or liver biopsy was recommended.
With this finding the patient referred to the gastroenterology center at King Abdul-Aziz Medical City in Riyadh for liver biopsy and possible ERCP.

Patient was seen there and started on ursodeoxycholic acid 300 bid with the differential diagnosis of sclerosing cholangitis versus Dubin-Johnson syndrome. In addition, to repeat MRCP after two years. However, later on March 2015 they decide to do a liver biopsy, which confirm the diagnosis of Dubin-Johnson syndrome.

Patient remain asymptomatic till November 2017 when he has been referred again to our clinic from the general practitioner with complain of right upper quadrant pain on and off associated with intermittent bleeding per rectum, occasional constipation and weight loss. During an examination, he was vitally stable, his abdomen was soft and lax no organomegaly or enlarged lymph nodes detected. Patient booked for upper and lower endoscopy.

UPPER ENDOSCOPY
- Esophagus Normal, Stomach Normal, Duodenum Normal.

Colonoscopy
- Scope passed up to Terminal ileum, a large polypoid tumor seen in rectum, 7-10 cm from anal verge looks malignant.

Then we requested CT scan chest / abdomen and pelvic which show:
- Eccentric polypoidal rectal thickening is seen involving the right lateral wall of the rectum with somewhat angulated margins, stranding of fat and thickening of the right mesorectal fascia and a questionable small lymph node in the right internal iliac chain. Further evaluation by MRI pelvis will be of help for better characterization of the lesion. No evidence of chest or abdominal metastasis is seen. No osseous metastasis is identified.

MRI Pelvis Show
- IMPRESSION: A low- mid rectal flat polypoidal soft tissue mass lesion is likely representing rectal cancer with suspicion of extramural venous invasion. No internal sphincter involvement is seen. No regional lymph node is seen.
- Rectum, Proctocolonoscopic biopsy show: moderately differentiated adenocarcinoma (low grade).

We break the bad news to our patient and he had referred to a near oncology center for further treatment.

Figure- a: HE staining x 100, b: HE staining x 200, c: HE staining x 400: malignant, cribriform and glandular proliferation with atypia, apoptotic cells and abnormal mitoses.


**DISCUSSION**

The molecular basis in Dubin-Johnson syndrome is absence or deficiency of human canalicular multispecific organic anion transporter MRP2/cMOAT caused by homozygous or compound heterozygous mutation(s) in ABCC2 located on chromosome 10q24 [10] this result in mainly in unconjugated hyperbilirubinaemia. There are reports describing the protective effect of bilirubin on the development of cancer emphasizing the antioxidative and anti-inflammatory effect of bilirubin [8]. In exploratory case-control study was based on 777 CRC patients and 986 controls from the Czech Republic, lower serum bilirubin levels were detected in CRC patients compared to the controls (p < 0.001); each 1 μmol/L decrease in serum bilirubin was associated with a 7% increase of CRC risk (p < 0.001) [6]. The fact that our patient with DJS developed colorectal adenocarcinoma few years after he diagnosed with DJS, This indicates that hyperbilirubinemia may not be sufficient to protect cancer development. Even more, there are case reports describe the coincidence of DJS and colon cancer [7]. Adenocarcinoma of the colon is the most common histopathological type of colorectal carcinoma. It ranks
fourth in men and third in women in Western Europe and the US and overall accounts for 98% of cancers of the large intestine [5]. Screening is recommended in many countries. In a recent study once only flexible sigmoidoscopy screening or flexible sigmoidoscopy and FOBT reduced colorectal cancer incidence and mortality on a population level compared with no screening. Screening was effective both in the 50- to 54-year and the 55- to 64-year age groups, absolute rate difference, 28.4 [95% CI, 12.1-44.7]; HR, 0.80 [95% CI, 0.70-0.92]) [4]. As it is widely considered a disease that affects people after the 5th decade of life, screening is not indicated before 50 years of age [9]. Until we find the answers to some of these doubts, doctors must put attention when evaluating a young adult with DJS and be aware of this risk and consequences of a late diagnosis of colorectal cancer.

CONCLUSION

In view of the all of the above, we think the link between DJS and colon cancer should be consider in future researches, to establish large epidemiological study will lead to understand more about the coincidence of DJS and colon cancer. Finding screening criteria in patient with DJS for colon cancer can help in early diagnosis and of course good outcome of the treatment.

Consent

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

REFERENCES

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