Young adult with liver cirrhosis, portal vein thrombosis, a reported case of α-1-anti-trypsin deficiency.

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Introduction

Alpha-1 antitrypsin is a protein with inhibitory capability over the proteolytic enzyme elastase. Since its first description in 1963, over 100 different α1 AT alleles have been described (Folch E., et al., 2007). The major clinical manifestations of a 1 AT deficiency relate to the function of a 1 AT and where it is made. A 1 AT serves as an inhibitor of neutrophil elastase (NE), a powerful, destructive proteolytic enzyme stored in neutrophils (Carrell R. W, et al., 1982) (Janoff A., 1985). The liver is the major site of a 1 AT gene expression, releasing 2 g of a 1 AT into the circulation daily. A 1 AT diffuses into most organs, where it protects extracellular structures from attack by NE released by activated or disintegrating neutrophils. The lower respiratory tract is particularly vulnerable to deficiency of a 1 AT, which normally represents > 90% of the anti-NE protective screen of the alveolar walls (Gadek J. E., et al., 1981)(Wewers, M. D., et al., 1987).

The presence of cirrhosis in α1-antitrypsin deficiency is low, approximately 2.2/100,000 for ZZ homozygotes. The male-to-female ratio was 2 to 1. In one-third of the patients alcohol could have been a co-adjuvant or aggravating factor in the liver disease (Folch E., et al., 2007). We describe a unique case of a 27 year-old man with a 1 AT, presented with liver cirrhosis portal vein thrombosis & multiple bony deformities.

Case presentation:
male patient 27 years old was admitted to tropical medicine department at (NHTMRI), works as a manual worker in printing factory, he lives and born in Manawat, Giza, Egypt, married 5 years ago and he is father of 1 daughter 2 years old, he used to smoke 20 cigarettes/day for 12 years with history of Hashish and alcohol intake. Positive contact to canal water and received anti-bilharzial treatment in the form of tablets 3 doses the last one is 10 years ago.

The main complaint of the patient is swelling of both lower limbs one year before admission. The condition started by gradual onset, slowly progressive course, the patient noticed swelling of both lower limbs up to knees, the condition is associated with fever, yellowish discoloration of the sclera and dark colored urine, the patient sought medical advice & was admitted to fever hospital, stay for 2 weeks then discharged after fever disappeared & jaundice subsided. Few months later the patient noticed increase of the lower limb oedema and was referred to NHTMRI, during hospital stay the patient developed an attack of haematensis, about 1 cup of fresh blood not followed by melena and stopped spontaneously. The patient gives past history of 2 orthopedic operations (correction of deformity). Another operation to repair tendon cut. There is positive consanguinity for his parents. His grandfather was diabetic.

By examination the followings are noticed: The vital signs of the patient are normal. There is tinge of jaundice. Congenital small bony swelling in the right mandible. Congenital maldeformity in the hands (the fingers of the left hand are thinner and taller than that of the right hand), 1st degree clumping of fingers. Fine tremors of both hands. Scar in the left index finger (tendon repair operation). Congenital maldeformity in both lower limbs (knock knees), scar of the orthopedic operation in the right thigh (correction of deformity). Both feet show marked pitting oedema. Right leg shows moderate pitting oedema up to knee. Left leg shows mild pitting oedema up to knee. Chest and heart are clinically free.

The upper border of the liver is encountered in the 5th intercostal space MCL, and the lower border is not felt. The lower border of the left lobe is felt 1 finger below xiphisternum with sharp border and firm consistency. The spleen & both kidneys are not felt; the Traub's area is resonant. No ascites is felt clinically. The laboratory and imaging studies reveal the followings:

Urine analysis: Alkaline, Protein: (+), Blood: (+), Bilirubin (+), Pus cells (3-4). R.B.Cs: (6-8). AST: 109 U/L (0-40), ALT: 59 U/L (0-40), T.BIL: 2.65 mg/dl, D.BIL: 0.1 mg/dl, Albumin: 2.3 g/dl, FBS: 96 mg/dl, Urea: 24 mg/dl, Creat.: 0.75 mg/dl, Uric acid: 3.5 mg/dl, CBC: HB: 12.5 g/dl, MCV: 87.6 fl, MCH: 29 pg, MCHC: 33.2 g/dl, WBCs: 3900, Gran.: 54%, Lymph: 39%, Platelets: 116,000, Comment: mild normocytic normochromic anemia, pancytopenia, E.S.R: 1st hour 25, INR: 1.39, AFP: 1.2 IU/ml, HBsAg: negative, HCV Ab: negative, ANA: negative, AMA: negative, ASMA: negative, S.Iron: 90 ug/dl (60 – 160) µg/dl, S.ferritin: 80 ug/L (20–300) µg/L, S.copper: 100 ug/dl (70-40) mg/dl, ceruloplasmin: 62 mg/dl (17.1–65.1) mg/dl, S.Homocysteine: 13.3 umol/l (5-15) umol/l, decreased activity of protein C activity: 50% (70-140)% , borderline protein S activity: 71% (70-140)%. Quantitative serum: Alpha-1 Antitrypsin: 1.37 g/l (0.9-2.09) g/l slit-lamp examination of the eyes shows no kayscer Fisheger ring. abdominal ultrasonography reveals Heterogenous liver with portal vein thrombosis. Splenomegaly with dilated lienorenal collaterals. No ascites.

Venous duplex on both lower limbs is normal. Chest x ray shows slight obliteration of the right costo-phrenic angle, cardiac shadow is normal. Spiral CT scan of the abdomen with contrast shows cirrhotic liver with no focal lesions, main portal vein thrombosis. Lienorenal collaterals, right sided pleural effusion, CT Chest shows bilateral apical emphysematous bullae & mild right sided pleural effusion.
Globules, fibrous septa, cirrhotic liver, portal vein thrombosis.

Upper GI endoscopy reveals esophageal varices grade I-II Incompetent cardia, then liver biopsy revealed a picture consistent with alpha-1-antitrypsin deficiency with early cirrhotic changes associated mild chronic cholestasis. Pulmonary function tests reveal mild restrictive lung disease. We start treatment with warfarin in small increasing dose aiming for portal recanalization and to prevent further thrombosis in other vessels.

Fig 1) CT scan of the abdomen showing cirrhotic liver, portal vein thrombosis.

Fig 2) CT chest showing apical emphysematous bullae.

Fig 3) prominent bridging fibrosis & occasional nodules-surrounded by fibrous septa

Fig 4) Interface hepatitis "piece meal necrosis"

Fig 5) mild cholestasis.
Globules

Fig 6) PAS positive-diastase resistant in periportal hepatocytes.
Discussion

Cases of alpha1–Antitrypsin deficiency may be sporadic cases but it should be in our differential diagnosis for screening of cases of liver cirrhosis with negative viral markers especially if it is associated with unexplained chest problems. This case is presented with portal vein thrombosis as a 1 AT is one of the major inhibitors of the intrinsic pathway of coagulation, concomitant decrease of protein C activity and low normal activity of protein S may pave the way for PV thrombosis in this case, on contrary there are published cases of a 1 AT deficiency associated with Henoch–Schönlein purpura (HSP) (Patterson CC, et al. 2005) (Meier P, et al. 2001).

Replacement therapy may not be of value in such a case as it will not lead to reversal of hepatic fibrosis, cessation of smoking is the first line management of this case, oral warfarin should be adjusted to obtain portal patency then liver transplantation should be considered, it is not only the curative treatment of the hepatic condition but also it will lead to stoppage of progression for the pulmonary complication like emphysema, the 5 year survival rate in these cases is excellent and may reach up to 95%, also screening of the family members is mandatory for early diagnosis and management.

References

حالة مسجلة لنقص إنزيم ألفا وان المضاد للتدريبسن وتسببه في حدوث جلطة بالوريد البابي وتليف بالكبد

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مقدمة:

منذ إكتشاف المرض عام 1963 ظهرت العديد من الطرازات الجينية للمرض وصلت حوالي 100 طراز. وحيث أن الأعراض المرضية لنقص أنزيم ألفا وان المضاد للتدريبسن مرتبطة بشكل وثيق مع الأعضاء التي يعمل بها هذا الأنزيم لذا فإنا غالبا ما نشاهد أعراضه المرضية في الكبد والرئتين.

الحالة:

تم رصد حالة شاب في العقد العشريني وهو يعاني من جلطة مزمنة بالوريد البابي الكبدي وبعد أجراء التحاليل وعمل أشعات عادية ومقطعية على البطن والصدر وعمل وظائف تنفس وعينة كبدية للمريض، ثبتت أصابعة هذا الشاب بمرض نقص أنزيم ألفا وان المضاد للتدريبسن.

النصوص:

ينبغي وضع مرض نقص أنزيم ألفا وان المضاد للتدريبسن في الحسبان أثناء البحث عن أسباب اعتلال الكبد بالتليف على الرغم من عدم شعوع المرض. تعتبر زراعة الكبد هي الحل الأمثل لهذا المريض من أجل إيقاف تدهور حالة الكبد والرئتين أيضاً.