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Congenital hepatic fibrosis in a young boy with congenital hypothyroidism: A case report

Fang-Fei Xiao, Yi-Zhong Wang, Fang Dong, Xiao-Lu Li, Ting Zhang

Abstract

BACKGROUND

Congenital hepatic fibrosis (CHF) is a rare autosomal recessive disorder characterized by variable degrees of periportal fibrosis and malformation of bile ducts. CHF is generally accompanied by a variety of conditions or syndromes with other organ involvement.

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A liver biopsy demonstrated congenital hepatic fibrosis. Four years later at the age of 26 he was readmitted, having had five episodes of haematemesis and melaena. Liver function tests were again normal (Table I). An oesophagogastrectomy Congenital hepatic fibrosis was performed but the patient collapsed and died on the second postoperative day.

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Congenital hepatic cysts or congenital hepatic fibrosis is seen in both Persian and Persian-cross cats as an autosomal dominant trait and in the Swiss Freiburger horse (also called the Franches-Montagnes horse) as an autosomal recessive trait that can be traced back to one stallion. Both are

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Congenital hepatic fibrosis, without renal cysts, may be the only feature of CDG type 1b, or phosphomannose isomerase deficiency (Fig. 9-11). 175, 180 The cases described by Pelletier and coworkers, 181 also shown by mutational analysis to be due to CDG type 1b, 182 revealed, in addition, dilation of the crypts of the large intestine (see Fig ...

Congenital hepatic fibrosis

Inherited Fibrocystic Liver Disease

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