

## The right hepatic artery syndrome

Kazumi Miyashita, Katsuya Shiraki, Takeshi Ito, Hiroki Taoka, Takeshi Nakano

Kazumi Miyashita, Katsuya Shiraki, Takeshi Ito, Takeshi Nakano, First Department of Internal Medicine, Mie University School of Medicine, Tsu, Mie 514-8507, Japan

Hiroki Taoka, First Department of Surgery, Mie University School of Medicine, Tsu, Mie 514-8507, Japan

Correspondence to: Katsuya Shiraki, MD, PhD, First Department of Internal Medicine, Mie University School of Medicine, 2-174 Edobashi, Tsu, Mie 514-8507, Japan. katsuyas@clin.medic.mie-u.ac.jp  
Telephone: +81-59-231-5015 Fax: +81-59-231-5201

Received: 2004-06-29 Accepted: 2004-08-12

### Abstract

Various benign and malignant conditions could cause biliary obstruction. Compression of extrahepatic bile duct (EBD) by right hepatic artery was reported as a right hepatic artery syndrome but all cases were compressed EBD from stomach side. Our case compressed from dorsum was not yet reported, so it was thought to be a very rare case. We present here the first case of bile duct obstruction due to the compression of EBD from dorsum by right hepatic artery.

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**Key words:** Right hepatic artery syndrome; Obstructive jaundice; Extrahepatic bile duct; Right hepatic artery; Endoscopic retrograde cholangiography; Cholangioenterostomy

Miyashita K, Shiraki K, Ito T, Taoka H, Nakano T. The right hepatic artery syndrome. *World J Gastroenterol* 2005; 11 (19): 3008-3009

<http://www.wjgnet.com/1007-9327/11/3008.asp>

### INTRODUCTION

In the biliary legion, anatomic variation is relatively common. Arterial anomalies are not infrequent findings during biliary surgery and variations in the positions of hepatic arteries<sup>[1-5]</sup> are well known. However, it has been rarely reported that the extrahepatic bile duct (EBD) is compressed by the vessels of hepatobiliary lesion.

### CASE REPORT

A 55-year-old Japanese man with an unremarkable medical and family history was admitted for fever and body weight loss. On laboratory examination, alkaline phosphatase was 1 403 IU/L (normal range: 120-340 IU/L) and gamma-glutamyl transpeptidase was 130 IU/L (normal range: 8-60 IU/L). An abdominal computed tomography

(CT) showed compression from the dorsum and stenosis of the EBD (Figure 1: thick arrow head, EBD; Figure 3) by the right hepatic artery (Figure 1: thin arrow head, anterior branch and posterior branch), but did not reveal a mass or lymph node swelling around the EBD. Endoscopic retrograde cholangiography (ERC) showed a stenotic lesion of the common hepatic duct (Figure 2). An intraductal ultrasonography and biopsy were performed transpapillary, but no malignant finding was reported. Drainage of the bile duct using an endoscopic nasobiliary drainage tube improved fever and normalized enzymes. The resultant diagnosis was cholangitis due to compression of the EBD by the right hepatic artery from the posterior side. As a treatment, the stenotic site of the common hepatic duct was removed and end-to-side hepaticojejunostomy was performed. The operation showed that the right hepatic artery had crossed EBD from posterior side of EBD and compressed it. Pathologic examination showed no malignancy or inflammatory cell invasion, but demonstrated sclerosis and severe tortuosity of right hepatic artery. The post-operative course was uneventful and the patient has been well for 1 year after the operation.



**Figure 1** An abdominal CT showed compression from the dorsum and stenosis of the EBD (thick arrow head: EBD) by right hepatic artery (thin arrow head: anterior branch and posterior branch), but did not reveal a mass or lymph node swelling around the EBD.



**Figure 2** 3D CT showed compression from the dorsum of the EBD (arrows).



**Figure 3** ERC showed a stenotic lesion of the common hepatic duct (arrow). An intraductal ultrasonography and biopsy were performed transpapillary, but no malignant finding was reported.

## DISCUSSION

Compression of the EBD by the right hepatic artery has been reported as a right hepatic artery syndrome, but in all six cases reported, the EBD was compressed from the stomach side. In particular in this patient, the right hepatic artery ran across the back of the bile duct, which is a standard variation. It is possible that the etiology of the obstructive jaundice in this patient was that the common hepatic artery

was short and proximal ramification of the right hepatic artery of the anterior segment branch and posterior segment branch or severe tortuosity of the right hepatic artery by arterial sclerosis.

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Science Editor Guo SY Language Editor Elsevier HK