Multiple acquired portosystemic shunts secondary to primary hypoplasia of the portal vein in a cat

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Internal Medicine

ABSTRACT. A 6-year 5-month-old spayed female Scottish Fold cat presented with a one-month history of gait abnormalities, increased salivation, and decreased activity. A blood test showed hyperammonemia and increased serum bile acids. Imaging tests revealed multiple shunt vessels indicating acquired portosystemic shunt. Histopathologic analysis of liver biopsy showed features consistent with liver hypoperfusion, such as a barely recognizable portal vein, increased numbers of small arterioles, and diffuse vacuolar degeneration of hepatocytes. These findings supported the diagnosis of primary hypoplasia of the portal vein/microvascular dysplasia, (PHPV/MVD). To our knowledge, this is the first case of feline PHPV/MVD that developed multiple acquired portosystemic shunts and presented with hepatic encephalopathy.

KEY WORDS: acquired portosystemic shunt, feline, hepatic encephalopathy, microvascular dysplasia, portal hypertension
the menace responses and the withdrawal reflex of the hind limbs were absent bilaterally, and the tactile placing response of the hind limb was decreased.

Complete blood count identified a mildly decreased hematocrit (29.2%). Abnormalities found on serum biochemical analyses were ALT 102 U/l (reference value, 22 to 84 U/l), BUN 11.9 mg/dl (17.6 to 32.8 mg/dl), CRE 0.7 mg/dl (0.8 to 1.8 mg/dl), NH₃ 325 µg/dl (23 to 78 µg/dl), IP 2.0 mg/dl (2.6 to 6.0 mg/dl), K 2.9 mEq/l (3.4 to 4.6 mEq/l). Fasting and postprandial serum bile acids were increased (115.1 and 58.6 µmol/l, respectively; reference value <10.0 µmol/l). Total protein, albumin, glucose, alkaline phosphatase, total bilirubin, calcium, sodium, chloride and thyroxine (T4) were within the reference ranges. The cat had tested negative for feline leukemia virus (FeLV) and feline immunodeficiency virus (FIV). Urinalysis showed no abnormalities. Abdominal ultrasonography identified multiple small, tortuous, aberrant vessels in cranial to the left kidney. No renal calculus or cystolith was observed. Other abdominal organs, including gall bladder, spleen, kidneys, adrenal glands, pancreas and intestines, were normal.

Helical computed tomography (CT) of the whole body was performed acquiring contiguous, 1-mm thick transverse images using an 80-slice multi-detector device with a pitch of 0.81 (Aquilion PRIME, Toshiba Medical Systems Co., Ltd., Tokyo, Japan). A single injection of iodinated contrast agent (Iohexol; Omnipaque, Daiichi Sankyo Co., Ltd., Tokyo, Japan) was administered intravenously (600 mgI/kg) and angiographic scans were acquired immediately, and 20, 40 and 120 sec post-injection. The CT examination confirmed multiple vascular abnormalities, including a tortuous vessel connecting the splenic vein with the left renal vein (Fig. 1A), multiple tiny and tortuous vessels close to the left kidney (Supplemental Movie 1), and connecting the splenic vein with the small vein around the urinary bladder (Fig. 1B and Supplemental Movie 2). Portal vein was dilated at the porta hepatis, and subsequent intrahepatic portal vein branches were thin and tortuous (Supplemental Movie 3). The liver parenchyma showed patchy enhancement.

Since the owner did not want further investigations, supportive therapy with lactulose (1 g/head; Monilac, Chugai Pharmaceutical Co., Ltd., Tokyo, Japan) orally twice a day, metronidazole (13 mg/kg; Flagyl, Shionogi & Co., Ltd., Osaka, Japan) orally twice a day, and a commercial therapeutic diet for hepatic health (Prescription Diet l/d Cat, Hill’s-Colgate Japan Ltd., Tokyo, Japan) were started.

Six months after the initial presentation, the cat returned for an exploratory laparotomy because of recurrent hyperammonemia and symptoms, including salivation and wandering. Coagulation testing was performed on citrated plasma. Prothrombin time was 11.0 sec (reference value, 8 to 11 sec), activated partial thromboplastin time was 53.4 sec (21 to 45 sec) and fibrinogen was 51 mg/dl (90 to 250 mg/dl). On gross examination, multiple small, tortuous blood vessels were noted in cranial to the left kidney. The liver was mottled in color, and the edge of the liver was rounded. No ascites was observed. Two liver biopsy specimens were taken from left and right medial lobe, and they were submitted for histopathological analysis.

Histopathologic evaluation revealed scarcely recognizable portal vein (Fig. 2A), increased numbers of small arterioles aberrantly located within the portal triad, mild fibrosis, and mild lymphocyte infiltration (Fig. 2B). Diffuse vacular degeneration of hepatocytes and mild sinusoidal congestion were present throughout the hepatic parenchyma. Taken together, this case was diagnosed as PHPV/MVD with multiple aPSS and hepatic encephalopathy.
The dosage of lactulose was increased to 2 g/head twice a day, and metronidazole and commercial therapeutic diet were continued for symptomatic therapy of hepatic encephalopathy. At the time of writing this article, 477 days since initial presentation, the cat is still alive. Her hepatic encephalopathy is well controlled and her quality of life is good.

To our knowledge, there are only two reports of feline PHPV/MVD without macroscopic PSS [6, 16]. The first case was a 9-month-old Birman that had episodes of ptyalism, apathy, and abnormal behavior (circling) with increased pre- and post-prandial bile acids and ammonia. A macroscopic shunt was not found on surgical abdominal exploration, and liver biopsy confirmed hepatic portovenous hypoplasia; therefore, PHPV was suspected [16]. The second case was a domestic shorthair cat that presented with a clinically suspected PSS and was diagnosed with MVD without a macrovascular PSS [6]. However, there were no detailed descriptions and prognosis reports of the cats with PHPV/MVD in these reports.

PHPV/MVD is commonly associated with cPSS in dogs [7]. In contrast, there are few reports of PHPV/MVD accompanied with cPSS in cats [6, 7]. At our institution, 30 cats were histologically diagnosed with PHPV/MVD by liver biopsy, from April 2008 to June 2017; in 26 cats, it was associated with cPSS, in two, it was associated with AV fistula, in one, there was no definitive diagnosis, and only one case of PHPV/MVD without macroscopic congenital anomalies (the present case) was observed (unpublished data). This result suggests that most PHPV/MVD cases are associated with cPSS in cats as well as in dogs.

Dogs with PHPV/MVD frequently have less severe clinical signs and a better long-term prognosis than do those with a PSS that are managed medically [4]. However, it is known that PHPV/MVD rarely forms multiple aPSS from portal hypertension and exhibits severe symptoms, such as hepatic encephalopathy. In a report of 42 dogs diagnosed with portal hypertension caused by PHPV/MVD, most of the dogs died naturally or were euthanized owing to the progression of the disease within the first few weeks [15]. In cats, it is unknown whether the formation of multiple aPSS indicates poor prognosis. In the present case, although formation of multiple aPSS was observed, the cat had good quality of life for 477 days, using supportive therapy for hepatic encephalopathy.

In the present case, the symptoms of hepatic encephalopathy occurred at middle age. The possible explanations are as follows: (1) congenital PHPV had been asymptomatic at young age but then symptomatic multiple aPSS with hepatic encephalopathy was developed at middle age; or (2) the owner was not aware of the symptoms although multiple aPSS with hepatic encephalopathy was developed.

One limitation is that portal vein pressure was not measured in this case. However, multiple portosystemic collateral vessels were observed in the present case, which strongly indicates portal hypertension. Multiple aPSS develop as a result of prehepatic or intrahepatic portal hypertension [3]. Contrary to cPSS, ligation of the multiple shunt vessels in aPSS is contraindicated. It is necessary to treat the underlying diseases. According to the previous 11 case reports of feline multiple aPSS, the cause were describes as follows: four cats had developed it after ligation of cPSS [8, 16, 17], two had portal vein thrombus [13, 14], two had congenital hepatic fibrosis [18], one had hepatic AV fistula [10], one had bile duct adenocarcinoma [9], one had chronic cholangitis [9], and four had no mention of the diagnosis [2, 11, 12]. Since liver cirrhosis is rare in cats, it is necessary to recognize that the underlying diseases in cats with multiple aPSS are different from those in dogs.

To our knowledge, the present case is the first feline case of multiple aPSS secondary to PHPV/MVD, suggesting that PHPV/MVD should be considered in the differential diagnosis of aPSS. Further epidemiological studies are required to characterize this rare disorder.

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