CASE REPORT

Successful endoscopic sclerotherapy with bile duct stenting for a vascular malformation neighboring the duodenal papilla in blue rubber bleb nevus syndrome

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Abstract

A 14-year-old girl, who had been diagnosed with blue rubber bleb nevus syndrome, was referred to our hospital because of iron deficiency anemia. Esophagogastroduodenoscopy revealed a dark and red-colored vascular malformation occurring just above the duodenal papilla. Because the lesion was regarded as the cause of the anemia, we performed polidocanol injection therapy with bile duct stenting. Since esophagogastroduodenoscopy performed a month later revealed a scarred ulcer, the bile duct stent was removed. She has been under observation as an outpatient without any symptoms.

KEYWORDS
blue rubber bleb nevus syndrome, duodenal papilla, endoscopic sclerotherapy, polidocanol, vascular malformation

INTRODUCTION

Blue rubber bleb nevus syndrome (BRBNS) is a rare systemic disease with multiple vascular malformations (VMs) of the skin, gastrointestinal tract, and other organs.¹ The skin lesions in BRBNS are often first noticed at birth or in the neonatal period, while those lesions rarely need interventions. In contrast, VM of the gastrointestinal tract can cause acute or chronic bleeding with consequent anemia.²

VMs of the gastrointestinal tract occur mostly in the small intestine and distal colon.³ To date, there have been some descriptions of small intestinal VMs in BRBNS, which were treated by endoscopic interventions.²,⁴–⁹ However, there are no reports of BRBNS with endoscopic treatment for VM near the duodenal papilla. Herein, we report a case of successful endoscopic sclerotherapy with bile duct stenting for a VM occurring just above the duodenal papilla in BRBNS.

CASE REPORT

A 14-year-old girl was referred to our hospital for a detailed examination of iron deficiency anemia. Her hemoglobin level was 3.5 g/dl at the time of the first visit. She was diagnosed with BRBNS at the age of 10 years due to the VMs in the skin, the cerebellum, the iris, and the duodenum. She also had a prior...
history of endoscopic sclerotherapy for duodenal VM. The details for the treatment at that time have been reported elsewhere.\(^1\) Her family history was unremarkable.

Esophagogastroduodenoscopy revealed a dark and red-colored VM occurring just above the duodenal papilla (Figure 1a). Capsule endoscopy revealed blood in the upper jejunum and the ileum. Double-balloon endoscopy revealed vascular malformations in the deep jejunum and ileum (Figure 1b,c). We first treated the VM in the jejunum and ileum by polidocanol injection therapy. Since the capsule endoscopy showed blood in the upper jejunum, we presumed that the duodenal lesion neighboring the papilla was also considered to be the cause of the anemia, and we decided to treat the lesion by endoscopy.

Magnetic resonance cholangiopancreatography showed neither dilatation nor stenosis of the bile duct, while the main pancreatic duct was thin under magnetic resonance cholangiopancreatography. To lessen the risk of cholangitis and to avoid iatrogenic damages in the pancreatic duct, we planned to perform polidocanol injection therapy under the bile duct stenting without pancreatic cannulation. Moreover, because the patient was young, we refrained from endoscopic sphincterotomy to preserve the function of the Oddi sphincter. After the guidewire was placed in the bile duct (Figure 2a), we punctured the center of the lesion and confirmed negative blood backflow. Consequently, we injected 1.5 ml polidocanol in total (Figure 2b). Subsequently, a bile duct stent (7-Fr, 7 cm, Flexima; Boston Scientific, Massachusetts, USA) was placed (Figure 2c; Supplementary video).

The patient complained of mild abdominal pain for 2 days after the treatment. Serum amylase was slightly elevated to 396 U/L on the day after treatment, but it returned to the normal level on the second day. During the course of the hospitalization, white blood cell count and C-reactive protein were not elevated. It was thus presumed that her abdominal pain was due to mild pancreatitis. Oral intake was started 3 days after the treatment. The patient was discharged 6 days thereafter.

Because esophagogastroduodenoscopy performed a month later revealed a scarred ulcer near the ampulla,
we removed the bile duct stent (Figure 3a, b). She has been under observation as an outpatient without any symptoms.

DISCUSSION

BRBNS is a rare disease characterized by systemic VMs. For VMs in the gastrointestinal tract, surgery, endoscopic treatment or medication can be applied, according to the location, size, and the number of lesions. However, in recent years, endoscopic treatments, especially sclerotherapy, have been increasingly reported due to their less invasiveness and simplicity.

An online search in PubMed of reports published during a period 2015–2021 identified seven cases of endoscopic treatment for duodenal and small intestinal VMs in BRBNS (Table 1).\(^2\)\(^4\)\(^5\) Of the cases and our present case, five patients were young females, and all the patients presented with symptoms of gastrointestinal bleeding. As for the location of the lesions, the small intestine was the more frequent site than the duodenum. Five of the eight patients were treated with sclerotherapy, two with endoscopic mucosal resection, and the remaining patient with Endoloop ligation. Our case is the first report of sclerotherapy for a VM near the duodenal papilla.

Polidocanol has been used for sclerotherapy of vascular lesions in the gastrointestinal tract. Recently, Igawa et al. reported that the optimal dose of polidocanol injection for small-bowel hemangioma is 0.2 ml per 1 mm of the lesion diameter.\(^10\) On the bases of this report, we injected 1.5 ml of polidocanol. Although mild abdominal pain was observed after the treatment, there were no serious complications during the clinical course. We believe that the recommended dosage seems to be appropriate, but further validation in a larger number of patients is warranted.

Before treatment, we discussed whether pancreatic duct stenting should be performed. However, because...
the pancreatic duct depicted by magnetic resonance cholangiopancreatography was very thin, we considered that the stent itself might pose a risk of pancreatitis. Fortunately, the patient did not develop severe pancreatitis after the treatment. However, the indication of pancreatic ductal stenting should be discussed in each case, and a practical and optimal management strategy of VMs at or around the duodenal ampulla should be established.

In conclusion, our experience suggests that polidocanol injection therapy is effective for the lesion near the duodenal papilla in patients with BRBNS. The necessity of bile and pancreatic stenting needs to be elucidated further.

CONFLICT OF INTEREST
Author Takayuki Matsumoto is the responsible and executive JGES member for DEN Open. The other authors declare no conflict of interest.

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SUPPORTING INFORMATION
Additional supporting information may be found in the online version of the article at the publisher’s website.

Supplement Video 1. Video image of polidocanol injection therapy. First, the guidewire was placed in the bile duct. Second, we punctured the center of the lesion and injected 1.5 ml polidocanol. Finally, a bile duct stent was placed.