Evaluation of Duodenal Angioectasia with Portal Hypertension

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Abstract

Background: A few studies have investigated duodenal lesions in patients with portal hypertension. Aim is to investigate duodenal angioectasia in patients with portal hypertension.

Methods: Sixty patients with duodenal angioectasia and portal hypertension were investigated between April 2009 and March 2012. The subjects were 29 males and 31 females ranging in age from 50 to 84 years (mean: 67.5). Endoscopic findings of duodenal angioectasia were investigated. We evaluated the therapeutic strategy for cases of bleeding duodenal angioectasia.

Result: The underlying pathologies of portal hypertension were liver cirrhosis in 56 patients, idiopathic portal hypertension in three patients and extrahepatic portal vein obstruction in one patient. Forty-one of the 60 patients had previously received endoscopic injection sclerotherapy for esophageal varices and the other nineteen patients had a coexistent high risk of esophageal varices. Gastric antral vascular ectasia was discerned in 29 cases. The location of the duodenal angioectasia was the duodenal bulb in 30 cases, descending portion in 13 cases and both the duodenal bulb and descending portion in 17 cases. Endoscopic findings of duodenal angioectasia were classified as follows: punctulate erythema (<1 mm), with or without oozing, and patchy erythema (a few mm), with or without oozing. Endoscopically, bleeding from the duodenal angioectasia was observed in 16 of 60 (26.7%) patients: punctulate erythema in 6 cases and patchy erythema in 10 cases. Bleeding from the patchy erythema type was discerned in 10 of 16 patients (62.5%). However, there was no bleeding in 43 cases of punctulate erythema involving the bulb. Argon plasma coagulation was successfully performed for 6 of 16 cases of bleeding duodenal angioectasia and the other 10 cases were followed-up with endoscopic observations.

Conclusions: Duodenal angioectasia in patients with portal hypertension is considered to be one of the lesions of portal hypertensive duodenopathy.

Keywords: Duodenal angioectasia; Portal hypertension; Portal hypertensive duodenopathy; Argon plasma coagulation

Introduction

Portal hypertension can result in either the reopening of collapsed embryonic channels or reversal of the flow within existing adult veins [1]. Whilst esophagogastric varices are the most common complication in patients with portal hypertension, ectopic varices defined by large portosystemic venous collaterals occurring anywhere in the gastrointestinal tract, other than the esophagogastric region, are less common and account for between 1% and 5% of all variceal bleeding [2,3]. Ectopic varices that are not esophagogastric are located predominantly in the duodenum, jejunum, ileum, colon, rectum, and enterostomy stoma. Bleeding from ectopic varices, which is rare in patients with portal hypertension is generally massive and life threatening. In addition, other mucosal lesions such as portal hypertensive gastropathy (PHG) [4-6], Portal Hypertensive Duodenopathy (PHD), And Portal Hypertensive Colopathy (PHC) may develop and cause bleeding in patients with portal hypertension.

A few studies have investigated duodenal lesions in patients with portal hypertension. Data on the frequency of PHD and clinical significances are scarce. Gupta et al. reported that PHG and PHD are distinct clinical and endoscopic entities [7]. Duodenal erosion in patients with portal hypertension have been reported to be one of the lesions of PHD [8].

The aim of this study is to assess duodenal angioectasia on endoscopy and its clinical characteristics in patients with portal hypertension.

Methods

This was a retrospective study of patients diagnosed with portal hypertension from 2009 to 2012 at the Sapporo Kosei Hospital. The study was designed to evaluate the assessment of duodenal angioectasia on endoscopy and its clinical characteristics in patients with portal hypertension.

Between April 2009 and March 2012, 60 patients with portal hypertension were diagnosed with duodenal angioectasia by routine upper endoscopy at Sapporo Kosei General Hospital. These 60 consecutive patients were studied (29 males and 31 females, ranging in age from 50 to 84 years [mean: 67.5]).

Endoscopic findings of duodenal angioectasia were classified as, punctulate erythema (<1 mm), with or without oozing (type-1a) (Figure 1A), or patchy erythema (a few mm), with or without oozing (type-1b) (Figure 1B) [9]. We evaluated the location and the endoscopic classification of duodenal angioectasia, with or without bleeding. The past therapeutic histories of esophageal varices and endoscopic findings of PHG and Gastric Antral Vascular Ectasia (GAVE) were reviewed. We evaluated the therapeutic strategy for cases of bleeding duodenal angioectasia.

The study was performed according to the tenets of the Declaration...
of Helsinki. Written informed consent was obtained from all patients prior to the procedures. The study was approved by the ethical committee of Sapporo Kosei Hospital.

Results

The underlying pathologies of portal hypertension were liver cirrhosis in 56 patients, idiopathic portal hypertension in 3 patients and extrahepatic portal vein obstruction in 1 patient. The liver functions of 56 cirrhotic patients, according to Child-Pugh classification, were: 25 class A, 22 class B and 9 class C. Forty-one of the 60 (68.3%) patients had previously received endoscopic injection sclerotherapy (EIS) for esophageal varices and the other nineteen patients had a coexistent high risk of esophageal varices. EIS had been performed on 11 of the 41 patients during the past year and no case of duodenal angioectasia was found. On the other hand, PHG was discerned in 25 of 60 (41.7%) patients. GAVE was discerned in 29 of 58 (50.0%) cases (except two post resection of the stomach) (Table 1).

The location of the duodenal angioectasia was the duodenal bulb in 30 cases, descending portion in 13 cases and both the duodenal bulb and descending portion in 17 cases. Duodenal angioectasia was discerned in the duodenal bulbs of 47 of the 60 (78.3%) patients. We evaluated the endoscopic findings of duodenal angioectasia and the locations were classified as follows: type-1a of the duodenal bulb in 26 cases, type-1b of the duodenal bulb in 4 cases, type-1a of the descending portion in 3 cases, type-1b of the descending portion in 10 cases, type-1a of the duodenal bulb and descending portion in 15 cases, and a combined type of type-1a of the duodenal bulb and type-1b of the descending portion in 2 cases.

Endoscopically, bleeding from the duodenal angioectasia was observed in 16 of the 60 (26.7%) patients, 3 of 30 in the duodenal bulb, 8 of 13 in the descending portion and 5 of 17 in the duodenal bulb and descending portion. In the 5 patients with involvement of the duodenal bulb and descending portion, bleeding was observed in the descending portion. Endoscopic classification of duodenal angioectasia and its bleeding rates was showed in Table 2. Bleeding was most common from the descending portion and in type 1b angioectasia. Next, we evaluated the relationship between the types of angioectasia and endoscopic findings. The bleeding frequency was as follows: in 0/43 (0%) of type-1a from the duodenal bulb, in 3/4 (75.0%) of type-1b from the duodenal bulb, in 6/18 (33.3%) of type-1a from the descending portion and in 7/12 (58.3%) of type-1b from the descending portion. There was no bleeding among 43 type-1a cases involving the bulb.

Endoscopic treatment was performed for 6 of 16 bleeding duodenal angioectasia (Figure 1C), and the other 10 cases were followed-up with endoscopic observations because of the amount of bleeding was small. Argon plasma coagulation (APC) in 5 of 6 cases, and APC plus endoscopic clipping in the remaining case, was performed successfully for these 6 bleeding patients.

Discussion

PHD was first reported as congestive gastrointestinal by
Thiruvengadam et al. [10] and its frequency varies, ranging from 8.4% to 51.7% of portal hypertensive patients [7,8,11,12]. Menchen et al. diagnosed PHD as a congestive vascular pattern of the duodenum and reported that it was found in 8.4% [13]. On the other hand, Barakat et al. found PHD in 51.4% of portal hypertensive patients, including erythema, erosions, ulcers, telangiectasia, exaggerated villous pattern, duodenal varices, and mixed lesions [14]. The hepatic reserve function was assessed by the Child-Pugh classification. In this study, there was no relationship between the grade with Child's grade A, B or C and duodenal angioectasia, similar to previous reports [7,11,13].

Findings of PHD have been observed more frequently in patients after EIS for esophageal varices [8,13] but several articles reported no direct association between EIS and an increased prevalence of PHD [7,12]. In this study, 41 of our 60 (68.3%) patients had previously received EIS for esophageal varices and the other nineteen patients had co-existing high risk esophageal varices. Duodenal angioectasia was diagnosed in 11 of 41 patients within one year of EIS and we suspect a relationship between EIS and the progression of duodenal angioectasia. This may imply that EIS cause artificial occlusion of the esophageal varices, leading to elevation of the portal pressure. El-Khayat et al. also has reported the worsening of PHG and PHE (including PHD) with presence of esophageal varices and after EIS. Duodenal angioectasia can cause occult bleeding in some cases and APC is a useful treatment for its bleeding. However, the sample size in this report is small and more investigations are necessary in larger numbers of patients by a long-term analysis.

References

Table 2: Endoscopic classification of duodenal angioectasia and its bleeding rates.

<table>
<thead>
<tr>
<th>Type</th>
<th>Duodenal bulb</th>
<th>Descending portion</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Type 1a</td>
<td>0/43(0%)</td>
<td>6/18(33.3%)</td>
<td>6/61(9.8%)</td>
</tr>
<tr>
<td>Type 1b</td>
<td>3/47(5.0%)</td>
<td>7/12(58.3%)</td>
<td>10/16(62.5%)</td>
</tr>
<tr>
<td>Total</td>
<td>3/47(6.4%)</td>
<td>13/30(43.3%)</td>
<td>16/77(20.8%)</td>
</tr>
</tbody>
</table>

Type 1a: punctulate erythema (< 1mm), with or without oozing
Type 1b: patchy erythema (a few mm), with or without oozing

The etiology of GAVE remains unknown, and it is thought to encompass a variety of conditions. The majority of GAVE patients have underlying cirrhosis and portal hypertension. Ikeda et al. reported that the development of GAVE associated with cirrhosis was dependent on the temporal progression of the disease [19]. On the other hand, Spahr et al. [20] reported that GAVE is not directly related to portal hypertension. Duodenal angioectasia and GAVE have a similar endoscopic appearance but the pathogenesis differs. In this study, GAVE was discerned in 50.0% of cases, however, of the 11 patients in whom duodenal angioectasia was not present before EIS, only two patients (20.0%) developed GAVE within one year of EIS. These findings may indicate no relationship between an elevation of portal pressure and GAVE.

Barakat et al. reported that PHD was documented as the cause of overt or occult bleeding in 9.5% of patients, and endoscopically, 10 (18.5%) of 54 patients had blood clots overlying the PHD and the episodes of overt bleeding were not severe. In this study, bleeding from the duodenal angioectasia was observed endoscopically in 16 of 60 (26.7%) patients. Bleeding was seen in 3 of 47 (6.4%) from the duodenal bulb and in 13 of 30 (43.3%) from the descending portion, and in 6 of 61 (9.8%) of type-1a and in 10 of 16 (62.5%) of type-1b. Bleeding was most common in the descending portion and in type 1b angioectasia. The bleeding was overt in 6 of 16 patients (37.5%), manifested as anemia and/or melena, and APC was successfully performed for these 6 patients. APC is a modality of non-contact electro coagulation that applies high frequency electronic energy into tissue to cause defined thermal effects, which can be used for thermal devitalization of the tissue as well as hemostasis [21]. The argon plasma seeks precisely the tissue surface with the lowest electrical impedance. Therefore, the tissue is coagulated to an even depth across the entire surface, and the danger of perforation is lower than with conventional coagulation procedures or YAG laser [22]. YAG laser has become the effective therapy for GAVE [23]. However, it has several disadvantages over APC. It has a higher set-up cost, and it is more bulky and more dangerous to perforate than APC. The depth of mucosal injury is limited with APC [22,24]. APC has been performed successfully to treat GAVE, and results in good temporary control of GAVE-associated hemorrhage [24-27]. In this study, APC was a safe and effective treatment for bleeding duodenal angioectasia.

The other 10 cases were followed up with endoscopic observations because of small amount of bleeding and this is appropriate for cases in which anemia is not worsening. Piccinini et al. diagnosed an atypical endoscopic picture of PHD and described an approach using APC [28].

On endoscopic examination, PHD shows various types, including redness, erosion, ulcers and telangiectasia, in the duodenal mucosa. In conclusion, duodenal angioectasia in patients with portal hypertension is considered to be one type of PHD, and its frequency increases with presence of esophageal varices and after EIS. Duodenal angioectasia can cause occult bleeding in some cases and APC is a useful treatment for its bleeding. However, the sample size in this report is small and more investigations are necessary in larger numbers of patients by a long-term analysis.

Acknowledgement
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