Superior Mesenteric Artery Syndrome in Patients with Crohn’s Disease: A Description of 2 Cases Studied with a Novel Magnetic Resonance Enterography (MRE) Procedure

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Case reports: We report 2 cases of Crohn’s disease patients with clinical suspicion of jejunal stricture who underwent MR-enterography with a novel approach. In fact, the examinations were performed including prone position of the patients inside the scanner, drinking of contrast medium during the examination, and prompt acquisition of fluoroscopic sequences. Both the exams showed an abrupt termination of the duodenum on its third portion and a decreased aortomesenteric distance, allowing the diagnosis of superior mesenteric artery syndrome.

Conclusions: A correlation between Crohn’s disease and superior mesenteric artery syndrome has never before been reported in the literature. The present study provides some practical steps that may be useful in order to improve MRE standard protocol in recognizing this condition while evaluating Crohn’s disease bowel lesions.

MeSH Keywords: Crohn Disease • Magnetic Resonance Imaging • Superior Mesenteric Artery Syndrome

Abbreviations: MRE – magnetic resonance enterography; SMA – Superior mesenteric artery; CD – Crohn’s disease; CDAI – Crohn’s disease activity index; CECT – Contrast-enhanced computed tomography

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Background

Superior mesenteric artery (SMA) syndrome, also called Wilkie’s syndrome, is an uncommon but well-recognized clinical entity characterized by compression of the third portion of the duodenum between the aorta and SMA, causing a chronic, intermittent or acute duodenal obstruction that can be complete or partial. SMA syndrome has mainly been described in 2 patient populations: patients who have undergone surgery, resulting in loss or distortion of the normal retroperitoneal fat (i.e., scoliosis surgery, aortic aneurysm repair, or bowel resection surgery) and patients who have had severe, rapid weight loss for any number of reasons [1]. Even if both these conditions are often concomitant in patients affected by Crohn’s disease (CD), a correlation between CD and SMA syndrome has never been reported.

In fact, according to randomized controlled trials, up to one-third of patients with CD currently undergo major abdominal surgery within 5 years [2]. In addition, weight loss is one of the most common signs of CD, representing a severe problem in patients who have undergone extensive surgical resections (e.g., short bowel syndrome) or with severe Crohn’s Disease Activity Index (CDAI).

Although CT-scan still represent the criterion standard imaging modality in diagnosing SMA syndrome, MR-enterography (MRE), which is a radiation-free imaging modality already included in the standard diagnostic algorithm of inflammatory bowel disease, could be helpful in recognizing the onset of SMA syndrome while evaluating intestinal Crohn’s disease lesions.

We report on 2 CD patients who underwent MRE for clinical suspicion of jejunal stricture. MRE, performed with a novel technical approach, demonstrated the finding of SMA syndrome, never reported in the Crohn’s literature up to now.

MRE – technique

Patients who undergo MRE of the small bowel can be examined by drinking 1500–2000 ml of a polyethylene glycol solution administered within 45 min before the beginning of the scan. On the other hand, intestinal distension can be obtained by injecting the contrast medium through a nasojejunal catheter (MR-enteroclysis). This latter procedure is more invasive and is often associated with patient discomfort, and it is actually considered optimal in case of jejunal stricture as it allows a better distension of the proximal intestinal segments.

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The patient is asked to lie in prone position inside the scanner and to drink the oral contrast agent through a drinking straw for the whole duration of MR-fluoroscopy (Figure 1).

MR-fluoroscopy is performed with several T2-weighted thick-slab turbo spin-echo sequences (TR/TE: $\times/900$ ms), usually employed in MR-cholangiopancreatography and MR-pyelography. Images are acquired on the coronal plane (section thickness: 100–180 mm) in order to include the gut and the stomach, and are obtained during and after the oral administration of contrast material, every 1 or 2 min (Figure 2).

When no strictured bowel tract is detected, the fluoroscopic-like images are then used as a guide for assessing the most adequate proximal bowel distension prior to beginning the standard MRE protocol. On the other hand, in case of abnormally and persistently dilated bowel segments followed by narrowed intestinal tracts, MR-fluoroscopic images are used as landmarks for targeting the following standard MRE sequences.

The standard protocol includes various T2-weighted pulse sequences along axial and coronal planes. High-resolution ultra-fast sequences, such as true-fast imaging with steady-state (True-Fisp; TR/TE: 4.20/2.10 ms, FA: 60°) and half-Fourier acquisition single-shot turbo-spin-echo (HASTE; TR/TE: $\times/80$ ms) with and without fat-suppression, are usually performed, together with diffusion-weighted imaging (DWI) sequences, using a diffusion factor $b$ fixed at 0, 400, and 800 s/mm$^2$. DWI helps to identify segments of bowel affected by CD and to assess disease activity [3].

When CD activity is challenging to assess, fat-suppressed three-dimensional gradient-echo images (THRIVE; TR/TE: minimum/minimum, FA: 10–15°) are added to the protocol either before and after intravenous gadolinium-based contrast material injection.

Figure 1. Picture showing the patient, in prone position inside the MR scanner, drinking oral contrast agent from a drinking straw during MR-fluoroscopy.
Case Report

Case 1

A 23-year-old female, with a diagnosis of CD A2L4bB2G0 according to the Paris classification [4], presented at our institution with an 8-month history of diarrhea and abdominal pain, associated with recurrent postprandial stabbing epigastric pain, nausea, and vomiting. She also stated a weight loss of 8 kg since the symptoms had begun, and she had been treated with adalimumab and azathioprine for the last 6 months but without relief of epigastric pain and nausea.

A MRE was performed with the clinical suspicion of proximal bowel obstruction. MR-fluoroscopy showed a persistently dilated duodenum with an abrupt termination at its third portion and mild gastric distention (Figure 3A). Conventional MRE sequences confirmed the presence of a grossly dilated duodenum abruptly narrowed under SMA and revealed an aortomesenteric distance of 4 mm (normal values: 10–28 mm) (Figure 3B, 3C). In addition, on delayed MRE sequence acquisition, obtained when the small bowel was sufficiently distended, wall-thickening of several bowel segments was also present (Figure 3D). Subsequently, a gastroduodenal endoscopy, performed to exclude the presence of strictures or ulcerations, revealed only a pulsatile compression of the third portion of the duodenum, further corroborating the diagnosis of SMA syndrome. The patient was managed with adalimumab and high-caloric enteral nutrition and gained 5 kg in 2 months, reporting a partial relief of symptoms.

Case 2

A 27-year-old female was first diagnosed with CD at age 16 years. Her phenotype was A1bL4bB2B3G1p according to the Paris classification [4]. She presented at our institution with a 10-month history of fatigue associated with intermittent postprandial dull epigastric pain, accompanied by intermittent non-bloody vomiting, early satiety, post-prandial fullness, and severe malnutrition (body mass index: 14.02 Kg/m²). She had a history of multiple bowel resections, with an estimated residual small-bowel length of 170 cm. She was originally on treatment with adalimumab and, after the worsening of symptoms, she had been managed with infliximab for the prior 12 months, without any clinical improvement. Physical examination revealed fullness in the epigastric region. After a gastroenterological consultation, she was suggested to undergo MRE to evaluate the presence of possible intestinal segments of narrowing.
MR-fluoroscopy showed dilated proximal duodenum with markedly slow passage of contrast beyond the third portion, with a cut-off in adjacency of SMA (Figure 4A). MRE also confirmed a persistently dilated duodenum prior to its passage under SMA. The aortomesenteric distance measured at the level of the third portion of the duodenum was markedly reduced (4 mm; normal values: 10–28 mm) (Figure 4B). In addition, on delayed MRE sequence acquisition, wall-thickening of some residual ileal segments was also present (Figure 4C).

The patient began treatment with metronidazole and prednisone and was managed with parenteral nutrition and gastric decompression via a nasogastric tube, followed by placement of a Dobbhoff catheter for post-pyloric feeding with Modulen-B.
1000 kcal/die (Nestlé, Lausanne, Switzerland). She tolerated oral diet 5 weeks later and started to gain weight (BMI: 15.62 Kg/m$^2$), with partial relief of symptoms.

**Discussion**

SMA syndrome was first described comprehensively by Wilkie. It is caused by the vascular compression of the third portion of the duodenum between the aorta and SMA [5].

Typically, patients present postprandial epigastric pain, nausea, and gastric distension. These symptoms are characteristically described as being partially relieved by lying prone or in the left lateral decubitus position, because these manoeuvres release tension on the small-bowel mesentery and thus on the aortomesenteric angle [6]. Identification can be a diagnostic dilemma and it is frequently made by exclusion, as SMA syndrome is rarely taken into account at first presentation of symptoms because of its low prevalence (0.013–0.3%) [7].

There is a slight female preponderance (64–66%), with 75% of cases occurring in individuals who range in age from 10 to 39 years [8, 9]. The incidence is higher in patients with severe weight loss (e.g., chronic debilitating disease or acute catabolic state) or history of surgery (e.g., scoliosis surgery, aortic aneurysm repair, or bowel resection surgery), as both these conditions may be responsible for a possible consequent distortion.
of the normal retroperitoneal fat with an increased mesenteric tension and caudal pull of SMA [7]. In fact, the duodenum is normally separated from the SMA and abdominal aorta by adipose tissue, which serves as a natural cushion to prevent extrinsic compression [8].

Although a direct association between SMA syndrome and CD has never been reported up to now, CD is a well-known condition that may cause abdominal pain, malabsorption of nutrients with weight loss, and intestinal obstruction, potentially occurring in any part of the digestive tract. In addition, according to Bougen and Peyrin-Biroulet, the rate of surgery at 5 years in patients affected by CD ranges between 25% and 33% [2]. Given the high risk of multiple bowel resection surgeries, it is not uncommon for these patients to experience short-bowel syndrome and severe weight loss. For this reason, close monitoring of CD patients is mandatory, as it can help identify patients at risk for disease relapses or complications [10].

Since the last decade, CD patients have benefitted from the advances of several imaging techniques, especially MRE. This technique provides data regarding the location, extent of bowel disease, and disease phenotype (i.e., fibrostenosing, inflammatory, or perforating) and is accurate for the detection of strictures, extra-luminal complications, and associated diseases. However, among complications of CD, SMA syndrome has never been described so far [11–13]. On the other hand, MRE has shown the advantage not only to depict anatomical abnormalities of the small-bowel but also to detect its functional disorders. In fact, the use of rapid dynamic thick-slab sequences (e.g., 70–180 mm), a technique also known as MR-fluoroscopy, allows instilling the oral contrast agent with a simultaneous MR-guide, until good distention of jejunal loops is obtained. Real-time MR-fluoroscopic studies of the small bowel have been adopted only in MR-enteroclysis so far. In this procedure, intestinal distention is achieved via administration of contrast material through a prepositioned nasojejunal balloon-tipped catheter, with serious procedure-related patient discomfort [14]. For this reason, the use of MR-enteroclysis has been limited in routine clinical practice, as opposed to the relatively discomfort-free MRE.

In our institution, we have recently introduced a novel procedure for performing MRE, which has mainly been employed in the clinical suspicion of proximal bowel stricture, with the aim to further improve the visualization of jejunal loops. This novel procedure requires the prone position of the patient inside the scanner and the drinking of the oral contrast medium through a drinking straw while acquiring the fluoroscopic sequences. This allows monitoring not only the small-bowel filling, but also the gastroduodenal emptying, providing information about the peristalsis efficiency and better depicting the proximal jejunal tract. In case of delayed emptying of an intestinal segment that is followed by a narrowed tract, the conventional MRE sequences are then focused on the region of interest.

Up to now, the radiological diagnosis of duodenal obstruction in SMA syndrome has been achieved by means of conventional contrast fluoroscopy, contrast-enhanced computed tomography (CECT), and angiography.

Conventional fluoroscopy usually demonstrates the dilatation of the proximal duodenum followed by a sharp narrowing as the duodenum travels underneath the SMA. On the other hand, CECT imaging has demonstrated that the 2 key signs of SMA syndrome are represented by an aortomesenteric distance of less than 8 mm and by an aortomesenteric angle of less than 22°. The former sign has a sensitivity and specificity of 100% for SMA syndrome, whereas the sensitivity and specificity of the latter are 42.8% and 100%, respectively [15].

In both our cases, MR-fluoroscopy showed a persistently dilated proximal duodenum with an abrupt narrowing at its third portion and with a mild gastric distention, findings consistent with what is seen on conventional contrast fluoroscopy in the SMA syndrome [15]. In addition, MRE revealed also the reduction of the aortomesenteric distance, the major sign of SMA syndrome.

However, MRE is a radiation-free exam whose role is well-established in the assessment and management of CD patients but not in evaluating SMA syndrome.

In fact, even if it allows an overall view of the abdominal cavity and the addition of fluoroscopy sequences can further improve its usefulness in highlighting the presence of an upper intestinal tract obstruction, measurement in the prone position may underestimate the aortomesenteric angle.

Nevertheless, consideration should be given to the fact that MRE could yield additional important information, with a high degree of accuracy, to confirm or dismiss the suspicion of SMA syndrome. Of course, the final diagnosis should be achieved through other imaging modalities (e.g., gastroscopy or CTA), which therefore would be performed in a wiser and more advisable way.

To the best of our knowledge, this is the first report in which the SMA syndrome has been detected by means of MR in a CD patient, and it is noteworthy to highlight the role of real-time MR-fluoroscopy in detecting this syndrome prior to performing MRE. In fact, it would have hardly been demonstrable either by means of conventional MRE (because distention of the duodenum and proximal jejunum is usually poor at the time of scanning) and MR-enteroclysis (because the nasojejunal catheter might go through the duodenal stricture). Moreover,
a clear distinction between an extrinsic compression and the presence of a duodenal or jejunal stricture is crucial, as this latter phenotype is strongly associated with an increased risk of relapse and complicated disease [16].

Both these latter points might represent important pitfalls to avoid when performing MR of the small bowel in CD patients, especially in those with clinically suspected proximal small-bowel disease.

Conclusions

In conclusion, either gastroenterologists and radiologists should be aware of the possible association of SMA syndrome with CD in order to reduce the symptoms and to prevent inappropriate or aggressive treatments. The prone position and the dynamic evaluation allowed by fluoroscopic sequences during simultaneous drinking of the oral contrast medium can be useful tools to add to the standard MRE protocol in order to recognize the presence of this pathological entity while evaluating CD small-bowel lesions.

Conflict of interest

None.

References:

1. Raman SP, Neyman EG, Horton KM et al: Superior mesenteric artery syndrome: Spectrum of CT findings with multiplanar reconstructions and 3-D imaging. Abdom Imaging, 2012; 37: 1079–88
2. Bouguen G, Peyrin-Biroulet L: Surgery for adult Crohn’s disease: What is the actual risk? Gut, 2011; 60: 1178–81
3. Morani AC, Smith EA, Ganeshan D, Dillman JR: Diffusion-weighted MRI in pediatric inflammatory bowel disease. Am J Roentgenol, 2015; 204: 1269–77
4. Levine A, Griffiths A, Markowitz J et al: Pediatric modification of the Montreal classification for inflammatory bowel disease: The Paris classification. Inflamm Bowel Dis, 2011; 17(6): 1314–21
5. Wilkie D: Chronic duodenal ileus. Am J Med Sci, 1927; 173: 643–49
6. Merrett ND, Wilson RB, Cosman, P, Blankin AV: Superior mesenteric artery syndrome: Diagnosis and treatment strategies. J Gastrointest Surg, 2009; 13: 287–92
7. Welsch T, Büchler MW, Kienle P: Recalling superior mesenteric artery syndrome. Dig Surg, 2007; 24: 149–56
8. Lee TH, Lee JS, Jo Y et al: Superior mesenteric artery syndrome: Where do we stand today? J Gastrointest Surg, 2012; 16: 2203–11
9. Blank V, Werlin S: Superior mesenteric artery syndrome in children: A 20-year experience. J Pediat Gastroenterol Nutr, 2006; 42: 522–25
10. Sauter B, Beglinger C, Girardin M et al: Monitoring disease activity and progression in Crohn’s disease. A swiss perspective on the IBD ahead ‘optimised monitoring’ recommendations. Digestion, 2014;89: 299–309
11. Horsthuis K, Bipat S, Bennink RJ, Stoker J: Inflammatory bowel disease diagnosed with US, MR, scintigraphy, and CT: Meta-analysis of prospective studies. Radiology, 2008; 247: 64–79
12. Mazzotti S, Blandino A, Scribano E et al: MR enterography findings in abdominopelvic extraintestinal complications of Crohn’s disease. J Magn Reson Imaging, 2013; 37: 1055–63
13. Mazzotti S, D’Angelo T, Rachchiusa S et al: Peritoneal inclusion cysts in patients affected by Crohn’s disease: Magnetic resonance enterography findings in a case series. Clin Imaging, 2016; 40: 152–55
14. Siddiki H, Fidler J: MR imaging of the small bowel in Crohn’s disease. Eur J Radiol, 2009; 69: 409–17
15. Fong JK, Poh AC, Tan AG, Taneja R: Imaging findings and clinical features of abdominal vascular compression syndromes. Am J Roentgenol, 2014; 203: 29–36
16. Flamant M, Trang C, Maillard O et al: The prevalence and outcome of jejunal lesions visualized by small bowel capsule endoscopy in Crohn’s disease. Inflamm Bowel Dis, 2013; 19: 1390–96