INTRODUCTION

Meckel’s diverticulum is regarded as one of the most common congenital anomalies of the gastrointestinal tract. Complications that arise due to a Meckel’s diverticulum include bleeding, obstruction, inflammation, and less commonly, perforation. Although complications from a Meckel’s diverticulum are rarely seen in the adult population, when observed, emergency surgery is often required. We report an interesting occurrence of a huge gangrenous Meckel’s diverticulum in an adult patient, which was managed with laparoscopic surgery.

CASE REPORT

A 45-year-old woman with no significant medical history and no prior surgical history presented with a one-day history of right-sided abdominal pain and vomiting. The pain was a constant, dull ache without any radiation. It was the patient’s first episode of abdominal pain that was associated with fever, chills and rigor. She did not have any change in bowel habits and there was no passage of blood per rectum. Physical examination demonstrated tenderness in the right iliac fossa with mild guarding. There was no palpable masses or abdominal distension. She had a raised leucocyte count of 13.6 × 10⁹/L, but other blood test investigations were unremarkable. The initial impression was that of an acute appendicitis or diverticulitis.

Computed tomography (CT) of the abdomen and pelvis revealed a 6.7 cm × 4.1 cm × 4.1 cm air-filled mass with dependent debris, suggestive of an infective lesion. The mass appeared to arise from the antimesenteric border of the small bowel, raising suspicion that it may be a Meckel’s diverticulum (Fig. 1). A diagnostic laparoscopy was performed in view of the patient’s clinical and CT findings. Intraoperatively, we found a large, distended and gangrenous Meckel’s diverticulum, measuring 6 cm × 4 cm, which was walled off by the surrounding omentum and small bowel (Fig. 2). The Meckel’s diverticulum and ileum were mobile towards the right iliac fossa trocar site. A 2.5 cm incision was made in the right iliac fossa, and the Meckel’s diverticulum with the ileum was brought through an Alexis extra-small wound protector extracorporeally (Fig. 3). A stapled resection of the Meckel’s diverticulum was then performed. The patient recovered well and was discharged on the third postoperative day. There were no postoperative complications such as bleeding, infection or formation of intra-abdominal abscess. Pathological results of the specimen confirmed the...
diagnosis of gangrenous Meckel’s diverticulum. The patient was followed up for a period of six months, and there was no recorded incidence of complications such as wound dehiscence, incisional hernia or postoperative adhesive intestinal obstruction.

**DISCUSSION**

Preoperative diagnosis of Meckel’s diverticulum is a challenge despite the availability of modern imaging. CT and technetium-99m radionuclide imaging are widely used, but the correct diagnosis is often established only at the time of laparotomy or laparoscopy. This is especially the case for Meckel’s diverticulum in adults, as symptoms are often nonspecific and complications are likely to occur at the time of presentation. Hence, a diagnosis of symptomatic Meckel’s diverticulum demands a high degree of suspicion, as an accurate preoperative clinical diagnosis is often difficult to make.

Although complications due to Meckel’s diverticulum are rare, they often present as surgical emergencies. These complications often warrant surgical resection of the diseased segment. A gangrenous Meckel’s diverticulum is very rare. There are only two previously reported cases of giant gangrenous Meckel’s diverticulum caused by axial torsion. In both cases, the patients underwent laparotomy, with one patient requiring resection of the involved small bowel with primary anastomosis. This highlights the difficulties encountered when the Meckel’s diverticulum becomes gangrenous, as it makes the surgery even more demanding.

As it was clinically difficult to distinguish between diverticulitis and appendicitis in our situation, laparoscopic evaluation was valuable in aiding diagnosis. Excision of the symptomatic Meckel’s diverticulum is considered the treatment of choice. With laparoscopy, excision has become minimally invasive, quicker, safer and more efficient. Although intracorporeal resection and anastomosis could have been performed, some of the considerations included spillage from enterotomy and the cost associated with the usage of laparoscopic staplers. In addition, the need for an eventual incision to extract the specimen from our patient made extracorporeal resection a more logical option. The use of a laparoscope allowed a much smaller incision, which upon resection of the diverticulum, enabled the rest of the small bowel to be easily examined.

The routine adoption of laparoscopy in complicated Meckel’s diverticulum is not encouraged. Careful patient selection and the laparoscopic experience of the surgeon are important considerations. However, we believe this case serves to highlight the safety and feasibility of performing a laparoscopic resection of a huge gangrenous Meckel’s diverticulum in an adult patient.

**REFERENCES**