Laparoscopic suture repair of idiopathic gastric perforation in Duchenne muscular dystrophy

Go Miyano, Hiroshi Nouso, Keiichi Morita, Hideaki Nakajima, Mariko Koyama, Masakatsu Kaneshiro, Hiromu Miyake, Masaya Yamoto, Koji Fukumoto, Naoto Urushihara

Access this article online Website: www.afrjpaedsurg.org DOI: 10.4103/0189-6725.170219 Quick Response Code:

ABSTRACT

We report herein an adolescent case of Duchenne muscular dystrophy (DMD) with idiopathic gastric perforation, in which emergency surgical repair was performed laparoscopically. A 14-year-old nonambulatory boy with DMD was brought to our emergency department with sudden onset of severe abdominal pain and distention. Plain radiograph and computed tomography confirmed the presence of free intraperitoneal air and intrapelvic effusion. The patient elected to undergo laparoscopic inspection with 4 trocars, revealing a focal perforation, 3-4 cm in diameter, on the upper gastric body near the diaphragm. The stomach was also found to have a thin wall without evidence of peptic ulcer disease or other abnormalities. An interrupted suture was placed using 4-0 PDS. The abdomen was extensively irrigated, and multiple J-Vac drains were left in situ. Total operation time was 90 min, and no intraoperative complications were encountered. Enteral feeding through a nasogastric tube was started on postoperative day 7. The postoperative course has been uneventful as of the 12-month follow-up. Pediatric surgeons should be aware of the increased risk of gastric perforation associated with DMD, and that laparoscopic repair can be safely performed even in emergency settings.

Key words: Duchenne muscular dystrophy, gastric perforation, laparoscopic repair

INTRODUCTION

Duchenne muscular dystrophy (DMD) is a fatal X-linked recessive disease, and the most common congenital neuromuscular disorder of children.^[1] Dystrophic

Department of Pediatric Surgery, Shizuoka Children's Hospital, Shizuoka, Japan

Address for correspondence:

Dr. Go Miyano, Department of Pediatric Surgery, Shizuoka Children's Hospital, 860 Urushiyama, Aoi-ku, Shizuoka 420-8660, Japan. E-mail: go1993@hotmail.co.jp changes in the smooth muscle of the gastrointestinal tract have been implicated as a cause of gastrointestinal dysfunction. [2] We report on an adolescent case of DMD with idiopathic gastric perforation, in which we performed emergency surgical repair laparoscopically.

CASE REPORT

A 14-year-old nonambulatory boy with DMD was brought to our emergency department with sudden onset of severe abdominal pain and distention. Complaints had been present for 5 h before he was transferred to our hospital, body temperature was 38.8°C, and he was tachycardic (heart rate, 130-150 beats/min). Systolic blood pressure was 70-80 mmHg. The associated risks of perioperative pulmonary dysfunction and cardiac failure in this patient were thus considered high. Plain radiography of the abdomen demonstrated free intraperitoneal air, and computed tomography also confirmed the presence of free intraperitoneal air and intrapelvic effusion [Figure 1]. The patient elected to undergo laparoscopic inspection to search for possible perforations in the digestive tract. Emergent inspection with 4 trocars, including one for suction and irrigation, revealed a focal perforation 3-4 cm in diameter on the fundus of the upper gastric body, near the diaphragm [Figure 2]. The stomach was also found to have a thin wall without evidence of peptic ulcer disease

This is an open access article distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 3.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as the author is credited and the new creations are licensed under the identical terms.

For reprints contact: reprints@medknow.com

Cite this article as: Miyano G, Nouso H, Morita K, Nakajima H, Koyama M, Kaneshiro M, *et al.* Laparoscopic suture repair of idiopathic gastric perforation in Duchenne muscular dystrophy. Afr J Paediatr Surg 2015;12:197-9.

or other abnormalities. An interrupted suture, using 4-0 PDS, was placed using an extra-corporeal knot-tying technique [Figure 3]. The abdomen was extensively irrigated, and 3 BLAKE drains were left *in situ*. Total operation time was 90 min, with only 3 ml of blood loss recorded, and no intraoperative complications were encountered. The early postoperative course was uneventful. A gastrografin study was performed on postoperative day 7, showing no leakage and complete closure of the gastric suture [Figure 4]. Enteral feeding through the nasogastric tube was then started. All drains were removed by postoperative day 9. The patient was discharged from hospital in a stable clinical condition on postoperative day 12, and as of the 12-month follow-up, remains alive and symptom-free.

DISCUSSION

The involvement of smooth muscle in addition to progressive dystrophic changes in striated muscle may cause clinical dysfunction of the gastrointestinal tract. [2,3]

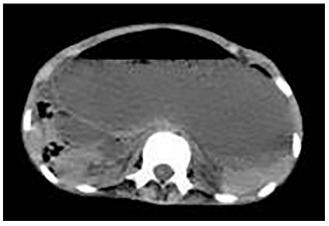


Figure 1: Preoperative abdominal computed tomography (CT) Abdominal CT confirm the presence of free intraperitoneal air and intrapelvic effusion

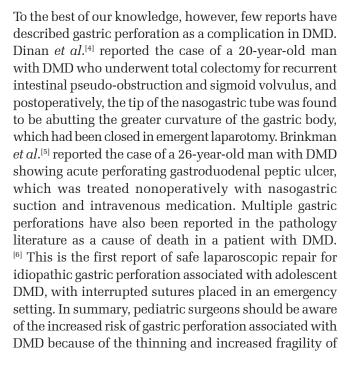




Figure 2: Laparoscopic view of the gastric perforation. Perforations (arrowheads) are present in the anterior wall of the upper stomach near the fundus



Figure 3: Laparoscopic repair of the perforation. The perforation is sutured laparoscopically using 4-0 PDS interrupted sutures



Figure 4: Postoperative upper gastrointestinal study. Complete closure of stomach perforation is confirmed, with no leakage apparent

the gastric wall, and that laparoscopic repair can be safely performed even in situations of emergency surgery.

Financial support and sponsorship

Conflicts of interest

There are no conflicts of interest.

REFERENCES

Emery AE. Population frequencies of inherited neuromuscular diseases — A world survey. Neuromuscul Disord 1991;1:19-29.

- Barohn RJ, Levine EJ, Olson JO, Mendell JR. Gastric hypomotility in Duchenne's muscular dystrophy. N Engl J Med 1988;319:15-8.
- Boland BJ, Silbert PL, Groover RV, Wollan PC, Silverstein MD. Skeletal, cardiac, and smooth muscle failure in Duchenne muscular dystrophy. Pediatr Neurol 1996;14:7-12.
- Dinan D, Levine MS, Gordon AR, Rubesin SE, Rombeau JL. Gastric wall weakening resulting in separate perforations in a patient with Duchenne's muscular dystrophy. AJR Am J Roentgenol 2003;181:807-8.
- Brinkman JM, Oddens JR, Van Royen BJ, Wever J, Olsman JG. Non-operative treatment for perforated gastro-duodenal peptic ulcer in Duchenne muscular dystrophy: A case report. BMC Surg
- Bevans M. Changes in the musculature of the gastrointestinal tract and in the myocardium in progressive muscular dystrophy. Arch Pathol (Chic) 1945;40:225-38.