Case Report

Delayed Presentation of Intussusception with Perforation after Splenectomy in Patient with Blunt Abdominal Trauma

Ibrahim Afifi, Hassan Al-Thani, Sajid Attique, Sherwan Khoschnau, Ayman El-Menyar, and Rifat Latifi

1 Department of Surgery, Section of Trauma Surgery, Hamad General Hospital, P.O. Box 3050, Doha, Qatar
2 Clinical Research, Section of Trauma Surgery, Hamad General Hospital, P.O. Box 3050, Doha, Qatar
3 Clinical Medicine, Weill Cornell Medical School, P.O. Box 24144, Doha, Qatar
4 Department of Surgery, Arizona University, Tucson, AZ 245005, USA

Correspondence should be addressed to Ibrahim Afifi; ibrahim_naeem@yahoo.co.uk

Received 16 September 2013; Accepted 25 November 2013

Academic Editors: A. Cho, F. Marchal, and A. R. Novotny

Copyright © 2013 Ibrahim Afifi et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Adult intussusception (AI) following blunt abdominal trauma (BAT) is a rare surgical condition. We present a case of delayed diagnosis of ileocecal junction intussusception with a perforation of small bowel in a 34-year-old male with a history of fall from height. Initial exploratory laparotomy revealed shattered spleen requiring splenectomy. Initial abdominal computerized tomography scanning (CT) scan showed dilated small bowel with no organic obstruction. Patient started to improve with partial distention and was shifted to rehabilitation unit. On the next day, he experienced severe abdominal distention and vomiting. Abdominal CT showed characteristic intussusception at the distal ileum. Secondary exploratory laparotomy revealed severe adhesions of stomach and small bowel to the anterior abdominal wall with dilated small bowel loops and intussusception near the ileocecal junction with perforation of small bowel. The affected area was resected and side-to-side stapled anastomosis was performed. Though small bowel intussusception is a rare event, BAT patients with delayed symptoms of bowel obstruction should be carefully evaluated for missed intussusception.

1. Introduction

Intussusception is the telescoping of one segment of the gastrointestinal tract into an adjacent one and usually occurs in children [1]. John Hunter was the first to describe the clinical and pathological characteristics of intussusceptions. Sir Fredrick Treves proposed the first management plan for intussusception which is being practiced to date [2]. Unlike children in whom most cases are idiopathic, the majority (80%) of adult intussusception (AI) cases have an underlying cause which could be due to development of tumors, fibrosis after surgery, and Meckel’s diverticula. Cases following blunt abdominal trauma are rare [3]. AI is relatively uncommon with different clinical presentation, diagnosis, and management compared to childhood intussusception [4]. AI is often diagnosed late during emergency laparotomy and the delayed diagnosis may be attributed to nonspecific presentation such as chronic colicky pain and intermittent partial intestinal obstruction associated with nausea and vomiting [5]. AI mainly needs surgical treatment which includes bowel resection with prior reduction of intussusception [4].

The increased utility of computerized tomography scanning (CT) helps in early evaluation of patients with abdominal pain and confirms the diagnosis for the possible AI without subsequent delay [1]. Definitive diagnosis of intussusception is possible due to its marked diagnostic appearance on CT imaging [6]. It appears as a complex soft-tissue mass, consisting of the outer intussuscipiens and the central intussusceptum. Though intussusception has a particular appearance on CT, it is difficult to identify the true etiological factors associated with its development [7]. Also, a high index of suspicion is required for diagnosing adult intussusception with nonspecific abdominal symptoms. However, such patients with history of prior episodes of partial intestinal
obstruction should undergo CT examination to rule out intussusception [6].

In the literature, small bowel intussusception due to blunt abdominal trauma (BAT) has not been extensively reported and only few case reports are described [8]. Herein, we present a case of ileocecal junction intussusception with perforation of small bowel following BAT in an adult male.

2. Case Report

A 34-year-old male patient presented with BAT with history of fall from 10-meter height. On presentation, the patient was hypotensive (blood pressure of BP 81/43 mmHg) with heart rate of 130 beat per min. He had depressed fracture of left frontoparietal skull bone, contusion with abrasion of left side of face, mild chest wall hematoma, and abdominal guarding. Patient was intubated immediately with rapid sequence induction. Focused abdominal sonography for trauma (FAST) was positive; chest X-ray showed left subcutaneous emphysema with fractures of the left 6th, 7th, and 10th ribs. Chest tube was inserted and massive transfusion protocol was activated. Initial laboratory investigations showed an INR of 1.2, low pH 7.16, base excess −9.4, low hematocrit (17%), high myoglobin (2549 ng/mL), creatinine 139 umol/L, low hemoglobin (10.1 g/dL), high white blood cell count (25.9 × 10³ u/L), normal platelet (246 × 10³ u/L), and high sensitive troponin T (90 ng/L). Patient was shifted to operation theatre immediately.

An exploratory laparotomy was performed which revealed shattered spleen requiring splenectomy; otherwise, formal exploration showed no other findings.

Postoperative laboratory investigation showed hemoglobin of 11.5 g/dL, WBC 13000 u/L, platelet 192000 u/L, creatinine 64 umol/L, myoglobin 1324 ng/L, and high sensitive troponin T 189 ng/L. Follow-up laboratory findings after 6 hours revealed low hemoglobin (8.5 g/dL), normal WBC (7300 u/L), and platelet count (130000 u/L).

A CT scanning of the head showed comminuted depressed fracture of left frontoparietal bone with minimal subdural hematoma, subarachnoid hemorrhage, tiny hemorrhagic contusions of left temporal area, and fracture of left zygomatic arch. Neck and chest CT scanning revealed evidence of C6 facet fracture and fractures of left 5–8 ribs anteriorly and left 6-7, 9–12 ribs posteriorly with left hemotorch and pneumothorax and bilateral lung contusions. With a smooth course in trauma intensive care unit (TICU), the patient was started enteral feeding through nasojejunal feeding tube after 24 hours and then he was extubated 2 days later. After removal of nasojejunal tube, the patient developed abdominal distention; for that nasogastric tube was inserted and 1100 mL fluid was removed initially. After a couple of days in the surgical ward, the patient showed a wax and wane course of abdominal distention. All serum electrolytes were within normal range.

Further CT scan of the abdomen showed dilated small bowel with no organic obstruction (Figure 1). Finally, he

![Figure 1: Abdominal CT scanning on day 9 of admission (day 5 after symptoms) showing peritoneal effusion, severely dilated small bowel, and suspected intussusception of the distal Ilium.](image)
started to improve with partial distention and the nasogastric tube was removed. A clear fluid was commenced and gradually advanced to full diet owing to the patient tolerance. Two days later, he experienced severe abdominal distention and vomiting, and therefore he was returned back to TICU. A clear fluid was started and the nasogastric tube was removed. A clear fluid was commenced and the nasogastric tube was removed. A clear fluid was commenced.

3. Discussion

Small bowel intussusception in BAT patients is uncommon and few cases were published. Historically, Leichtenstern reported a series of 326 cases of intussusception; of these cases, only 8% had past history of trauma as an etiology for intussusception [9]. The mechanism of traumatic intussusception is not well defined yet. Komadina and Smrkolj [10] described that the generation of localized bowel spasm following abdominal trauma, abnormal peristalsis, and intramural hematoma might be the possible mechanism for the development of small bowel intussusception. Trauma causes adrenergic stimulation of the sympathetic nerves of the gastrointestinal tract with a resultant spasm of the sphincters and segmental spasm of bowel. The direct blunt abdominal trauma causes positive wave pressure which gives rise to enough increased peritoneal cavity pressure to drive many spastic areas into adjacent areas of relative dilatation. Other factors that can be added to the development of intussusception include recently ingested meal, severe fright, and the possible existence of autonomic nerve imbalance in cases of severe stress of trauma. Yet, it is apparent that without trauma as the trigger mechanism, these factors never would have produced remarkable lesions.

Duncan and colleagues [11] identified 6 cases of intussusception as a cause of intestinal obstruction after laparotomy among abdominal trauma patients. Five cases were diagnosed with jejunojejunal intussusception; one had jejunoileal and one had ileocolic intussusception. Manual reduction was successfully used to treat all patients and recurrence was not observed in any of cases [11]. Bashir and Lynch [12] reported a case of double jejunojejunal intussusception causing intestinal obstruction after blunt abdominal trauma. The patient recovered completely with operative manual reduction. Berne et al. [13] reported a case of double intussusception after laparotomy for penetrating liver injury. The authors suggested that intussusception can be reduced manually without the need for surgical resection.

A recent case report described an ileoileal intussusception after BAT in patient presented with symptoms of intestinal obstruction after a fall from a ladder [14]. Though the authors resected the necrotic bowel segment, they have advocated the manual reduction to overcome the possible complications of surgical resection.

Benjelloun et al. [3] presented another case of ileoileal intussusception with Meckel's diverticulum. Meckel's diverticulum was a lead point for intussusception in BAT patients, and trauma with disturbed bowel motility was the unlatching mechanism of intussusception [3]. The authors performed a diverticulectomy with small bowel resection to attain successful outcome. Another BAT case presented after motor vehicle crash and showed jejunal intussusception on abdominal exploration. The patient was managed by manual reduction with unremarkable postoperative course [8].

Controversial findings have been reported for the exact treatment of adult intussusception. However, surgical resection has been advocated by many investigators [15–17]. However, Kitamura et al. [18] suggested simple reduction for posttraumatic and idiopathic intussusceptions where no
obvious pathological cause is suspected in the bowel [4]. In our case, surgical resection was used to manage the intussusception at ileocecal junction. The affected area was resected using staplers and side-to-side anastomosis.

In conclusion, the delayed presentation of small bowel intussusception after BAT is uncommon phenomenon. The diagnosis is difficult due to nonspecific and episodic symptoms which require a high index of suspicion. Abdominal CT is the modality of choice for diagnosis. Treatment requires resection of the small bowel if it is nonviable. Therefore, BAT patients with symptoms of bowel obstruction should be carefully evaluated for the possibility of intussusception.

Acknowledgments

The authors thank all the trauma surgery staff for their cooperation. The authors have no conflict of interests and no financial issues to disclose. All authors read and approved the case report. This case report has been approved by the Medical Research Center (IRB no. 13310/13), Hamad Medical Corporation, Doha, Qatar.

References

[1] G. Gayer, S. Apter, C. Hofmann et al., “Intussusception in adults: CT diagnosis,” Clinical Radiology, vol. 53, no. 1, pp. 53–57, 1998.
[2] M. D. Stringer and L. E. Willetts, “John Hunter, Frederick Treves and intussusception,” Annals of the Royal College of Surgeons of England, vol. 82, no. 1, pp. 18–23, 2000.
[3] E. B. Benjelloun, A. Ousadden, K. Ibnmajdoub, K. Mazaz, and K. A. Taleb, “Small bowel intussusception with the Meckel's diverticulum after blunt abdominal trauma: a case report,” World Journal of Emergency Surgery, vol. 4, no. 1, article no. 18, 2009.
[4] S. Yalamarthi and R. C. Smith, “Adult intussusception: case reports and review of literature,” Postgraduate Medical Journal, vol. 81, no. 953, pp. 174–177, 2005.
[5] R. S. Gordon, K. B. O'Dell, A. J. Namon, and L. B. Becker, “Intussusception in the adult—a rare disease,” Journal of Emergency Medicine, vol. 9, no. 5, pp. 337–342, 1991.
[6] G. Gayer, R. Zissin, S. Apter, M. Papa, and M. Hertz, “Adult intussusception—a CT diagnosis,” British Journal of Radiology, vol. 75, no. 890, pp. 185–190, 2002.
[7] D. G. Begos, A. Sandor, and I. M. Modlin, “The diagnosis and management of adult intussusception,” American Journal of Surgery, vol. 173, no. 2, pp. 88–94, 1997.
[8] Z. T. Stockinger and N. McSwain Jr., “Intussusception caused by abdominal trauma: case report and review of 91 cases reported in the literature,” Journal of Trauma, vol. 58, no. 1, pp. 187–188, 2005.
[9] H. Falor, “Multiple intussusceptions, direct and retrograde, of traumatic origin,” Annals of Surgery, vol. 127, no. 4, pp. 730–737, 1948.
[10] R. Komadina and V. Smrkoli, “Intussusception after blunt abdominal trauma,” Journal of Trauma, vol. 45, no. 3, pp. 615–616, 1998.
[11] A. Duncan, T. F. Phillips, S. J. Sclafani et al., “Intussusception following abdominal trauma,” Journal of Trauma, vol. 27, no. 11, pp. 1193–1199, 1987.