Case Report

Gastric Trichobezoar: An Enduring Intrigue

S. Mewa Kinoo and B. Singh

Department of Surgery, Nelson R Mandela School of Medicine, University of KwaZulu-Natal, 719 Umbilo Road, Congella 4013, South Africa

Correspondence should be addressed to S. Mewa Kinoo, smewakinoo@gmail.com

Received 26 September 2012; Accepted 8 November 2012

1. Introduction

A bezoar is a mass of undigested material within the gastrointestinal tract. The term bezoar derives from the Arabic word Badzehr, which means antidote [1]. Bezoars were used as antidotes against plague, snake-bite, leprosy, and epilepsy by physicians from 12th to 18th century [2]. Trichobezoar is from the Greek word trich which means hair [3]. A trichobezoar is a mass of undigested hair within the gastrointestinal tract. Trichobezoars are often associated with trichotillomania (hair pulling), and trichophagia (hair swallowing). Trichotillomania may be unconsciously or unintentionally done and is part of the DSM IV psychiatric classification of impulse control disorders [4, 5]. In up to 18% of patients with trichotillomania, trichophagia occurs; one third of patients with trichophagia develop trichobezoars [6]. Trichobezoars most commonly occur in adolescent females [7]. The site of hair pulling is most commonly from the scalp, but can occur from the eyelashes, eyebrows, and pubic area [8].

2. Case Presentation

A 16-year-old female was referred to our surgical services with a problem of epigastric pain associated with an epigastric mass. The patient resided in a local children’s home due to poor social circumstances. On questioning, patient reported nonspecific pain of approximately 3 months duration with no dyspeptic symptoms; the patient reported no early satiety and no history of weight loss. Examination revealed a well looking girl, with a nondistended abdomen. Palpation of her abdomen revealed a large, firm mobile epigastric mass that was minimally tender. Ultrasound and Computed Tomography (CT) scan of the abdomen both confirmed the presence of a large gastric mass with internal air loculi involving the entire stomach with extension into the duodenum but not into the jejunum (Figure 1). At endoscopy a trichobezoar, involving almost the entire capacity of the stomach, extending from the distal oesophagus into the duodenum was noted (Figure 2). Prompted by these diagnostic findings, on further enquiry, the patient admitted to trichophagia. The patient was referred to the hospital psychiatrist, social worker, and psychologist for counselling. Due to the size of the trichobezoar and potential for complications, operative removal of the trichobezoar was undertaken successfully via a gastrostomy (Figure 3). The patient was discharged well on day 5 into the care of the psychologists.

3. Discussion

Trichobezoars commonly occur in adolescent females, often with an underlying psychiatric or social problem. Clinical presentation of these patients may be confusing as often they
are not forthcoming with a history of trichophagia either due to embarrassment or the unintentional nature of the problem. Although this is a rare condition, numerous case reports and series have been reported as high mortality may follow complications associated with this condition.

Trichobezoars in humans were first described from a post mortem by Swain in 1854 [9]. The postulated reason for formation in the stomach is that hair is undigestable and due to its smooth nature cannot be propelled with peristalsis and over time forms a bezoar within the stomach. This bezoar can extend distally from the stomach into the caecum. Extension of the bezoar from the stomach into the jejunum or further on is referred to as “Rapunzel syndrome,” first described by Vaughan Jr. et al. in 1968 [10]. Rapunzel was a long haired girl in a German fairy tale by Grimm brothers. Bezoars can also be found distally in the gastrointestinal tract without continuity with the stomach bezoar due to breakage and distal propulsion. Trichobezoars continue to grow in size with continued ingestion of hair and this increases the risk of severe complications. The most common of these complications that have been reported over the years include gastric mucosal erosion, ulceration, and perforation of the stomach or the small intestine, gastric outlet obstruction, intussusception, obstructive jaundice, protein-losing enteropathy, pancreatitis, and death [11–16].

Presentation ranges from nonspecific abdominal or epigastric pain, to a range of complications as mentioned. Clinical examination often reveals a large mobile epigastric mass that may be indentable, the so-called Lamerton’s sign [17]. Endoscopy is usually diagnostic. The hair appears black (despite the normal hair colour) due to denaturing of the hair protein by the acid. The most common diagnostic tool used in the literature is a CT scan, with a typical image showing a well-defined intraluminal ovoid heterogeneous mass with interspersed gas [18, 19].

Management options include endoscopic removal, laparoscopic removal, or via laparotomy. Gorter et al., in a retrospective review of 108 cases of trichobezoar, evaluated the available management options [20]; it was noted that whereas 5% of attempted endoscopic removals were successful, 75% of attempted laparoscopies were successful. However, laparotomy was 100% successful and thus favoured as their management of choice.

4. Conclusion

Trichobezoars should be considered as a differential diagnosis in a young female patient with a mobile epigastric mass. Diagnosis can be easily made with the use of CT scan and endoscopy. Management almost always requires surgical removal. It is emphasized that the majority of these patients have an underlying psychiatric or social disorder. A multidisciplinary approach is essential to prevent recurrence of the problem.

References

[1] A. Samad, M. Ahmad, and Z. Latif, “Bezoars: a review and report of two cases,” Journal of the College of Physicians and Surgeons Pakistan, vol. 7, no. 6, pp. 263–265, 1997.
[2] S. Khattak and K. Asghar, “Trichobezoar,” Gomal Journal of Medical Sciences, vol. 2, no. 1, pp. 25–26, 2004.
[3] M. E. Rabie, A. R. Arishi, A. Khan, H. Ageely, G. A. S. El-Nasr, and M. Fagihi, “Rapunzel syndrome: the unsuspected culprit,” World Journal of Gastroenterology, vol. 14, no. 7, pp. 1141–1143, 2008.

[4] American Psychiatric Association, Diagnostic and Statistical Manual of Mental Disorders, American Psychiatric Association, Washington, DC, USA, 4th edition, 2000.

[5] G. A. Christenson and S. J. Crow, “The characterization and treatment of Trichotillomania,” Journal of Clinical Psychiatry, vol. 57, no. 8, pp. 42–49, 1996.

[6] C. Bouwer and D. J. Stein, “Trichobezoars in trichotillomania: case report and literature overview,” Psychosomatic Medicine, vol. 60, no. 5, pp. 658–660, 1998.

[7] M. R. Phillips, S. Zaheer, and G. T. Drugas, “Gastric trichobezoar: case report and literature review,” Mayo Clinic Proceedings, vol. 73, no. 7, pp. 653–656, 1998.

[8] G. A. Christenson, T. B. Mackenzie, and J. E. Mitchell, “Characteristics of 60 adult chronic hair pullers,” American Journal of Psychiatry, vol. 148, no. 3, pp. 365–370, 1991.

[9] E. M. Chisholm, H. T. Leong, S. C. S. Chung, and A. K. C. Li, “Phytobezoar: an uncommon cause of small bowel obstruction,” Annals of the Royal College of Surgeons of England, vol. 74, no. 5, pp. 342–344, 1992.

[10] E. D. Vaughan Jr., J. L. Sawyers, and H. W. Scott Jr., “The Rapunzel syndrome. An unusual complication of intestinal bezoar,” Surgery, vol. 63, no. 2, pp. 339–343, 1968.

[11] D. E. Ventura, F. A. Mardiros Herbell, S. T. Schettini, and C. Delmonte, “Rapunzel syndrome with a fatal outcome in a neglected child,” Journal of Pediatric Surgery, vol. 40, no. 10, pp. 1665–1667, 2005.

[12] M. H. Mehta and R. V. Patel, “Intussusception and intestinal perforations caused by multiple trichobezoars,” Journal of Pediatric Surgery, vol. 27, no. 9, pp. 1234–1235, 1992.

[13] H. Schreiber and H. C. Filston, “Obstructive jaundice due to gastric trichobezoar,” Journal of Pediatric Surgery, vol. 11, no. 1, pp. 103–104, 1976.

[14] A. Hossenbocus and D. G. Colin Jones, “Trichobezoar, gastric polyposis, protein losing enteropathy and steatorrhoea,” Gut, vol. 14, no. 9, pp. 730–732, 1973.

[15] R. N. Shawis and C. M. Doig, “Gastric trichobezoar associated with transient pancreatitis,” Archives of Disease in Childhood, vol. 59, no. 10, pp. 994–995, 1984.

[16] A. R. Jensen, C. T. Trankiem, S. Lebovitch, and H. Grewal, “Gastric outlet obstruction secondary to a large trichobezoar,” Journal of Pediatric Surgery, vol. 40, no. 8, pp. 1364–1365, 2005.

[17] A. J. Lamerton, “Trichobezoar: two case reports—a new physical sign,” American Journal of Gastroenterology, vol. 79, no. 5, pp. 354–356, 1984.

[18] J. Tamminen and D. Rosenfeld, “CT diagnosis of a gastric trichobezoar,” Computerized Medical Imaging and Graphics, vol. 12, no. 6, pp. 339–341, 1988.

[19] F. Navab and J. Sabol, “Images in clinical medicine,” New England Journal of Medicine, vol. 336, no. 24, p. 1721, 1997.

[20] R. R. Gorter, C. M. F. Kneepkens, E. C. J. L. Mattens, D. C. Aronson, and H. A. Heij, “Management of trichobezoar: case report and literature review,” Pediatric Surgery International, vol. 26, no. 5, pp. 457–463, 2010.