



A Rare Entity of Meckel's Diverticulum at Mesenteric Location : Two Case Reports

Sopan N. Jatal^{a++*}, Sudhir Jatal^{b#} and Sachin Ingle^{ct†}

^a MIMSR Medical College, Latur, Jatal Hospital and Research Centre, Latur, India.

^b Jatal Hospital and Research Centre, Latur, Tata Hospital, Mumbai, India.

^c Department of Pathology, MIMSR Medical College, Latur, India.

Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

Article Information

Open Peer Review History:

This journal follows the Advanced Open Peer Review policy. Identity of the Reviewers, Editor(s) and additional Reviewers, peer review comments, different versions of the manuscript, comments of the editors, etc are available here: <https://www.sdiarticle5.com/review-history/100552>

Case Report

Received: 24/03/2023

Accepted: 26/05/2023

Published: 27/05/2023

ABSTRACT

A Meckel's Diverticulum is the most common congenital anomaly of the gastrointestinal tract. Meckel's Diverticulum is usually located at the antimesenteric border of the small intestine and it is one of the cardinal findings of Meckel's Diverticulum. The Mesenteric location is an unusual site of Meckel's Diverticulum.

The etiology of the anomaly of Mesenteric Meckel's Diverticulum was due to the persistence of a short vitelline artery that creates a Mesodiverticular band from mesentery to the tip the diverticulum, which diverts the diverticulum away from the antimesenteric border to Mesenteric location during the elongation and growing process. We came across such two cases of mesenteric Meckel's Diverticulum (MMD). After each and every case of appendectomy or laparotomy, we traced 2 feet of small intestine (ileum) and to our surprise, we noticed such rare

⁺⁺ Ex. Professor;

[#] Fellow in Colorectal Cancer;

[†] Professor;

*Corresponding author: E-mail: jatalhospital@gmail.com;

and uncommon location of Mesenteric Meckel's Diverticulum, in two cases admitted in our hospital with classical signs and symptoms of acute appendicitis but after appendectomy we searched for Meckel's diverticulum as per our routine surgical procedure. Herein we are presenting two such rare and interesting cases with Mesenteric location.

Keywords: Case report; Meckel's diverticulum; mesenteric location; resection and anastomosis.

1. INTRODUCTION

"The omphalomesenteric duct persists forming on out pounding diverticulum called Meckel's Diverticulum, which is the most common congenital anomaly of the gastrointestinal tract". [1]

"The classical location of Meckel's Diverticulum is at the antimesenteric border of the bowel. The location of Meckel's Diverticulum in the opposite mesenteric side is extremely rare" [2,3].

"From 1941 to December 2012 only 32 cases of mesenteric Meckel's Diverticulum reported in the literature" [4,5].

"Etiology of the anomaly was due to the 10% persistence of a short vitelline artery that creates a Mesodiverticular band from the mesentery to the tip of the Meckel's Diverticulum which diverts the diverticulum away from the antimesenteric border during the elongation and growing process. Short Mesodiverticular band extending from the mesentery to the tip of the Meckel's Diverticulum, acts as Hamstring action and pulling the Meckel's Diverticulum away from the antimesenteric border of small intestine, which is classically seen in our two cases" [3,5].

"The Mesenteric location of Meckel's Diverticulum may erode mesentery and rupture

in to the mesenteric vasculature during the inflammatory process that may cause severe bleeding. Therefore, the surgical decision should be standard resection and anastomosis, even if the lesion is incidentally detected, during appendectomy or laparotomy" [4,5].

2. CASE PRESENTATION

Case I

A 17 years' adolescent boy in the year 1995, admitted in our center with complaints of pain in right lower abdomen, vomiting for 2 days. Abdominal examination revealed tenderness at McBurney's point. Haemotological investigation show leukocytosis. Ultrasound of abdomen revealed probe tenderness in right iliac fossa, considering the clinical and ultrasonography findings the diagnosis of Acute appendicitis was made. So the abdomen was opened by McBurney's incision, on exploration revealed Acute appendicitis and appendectomy was performed. Routinely traced the terminal 2 feet of ileum proximal to ileo-caecal junction. A diverticulum was found at the Mesenteric border of the small bowel of size 4x2 cm in length (Fig. 1a,1b). The patient recovery uneventful and patient discharged 7th postoperative day. Histopathological report of the specimen shows Meckel's Diverticulitis with no heterotopic tissue.

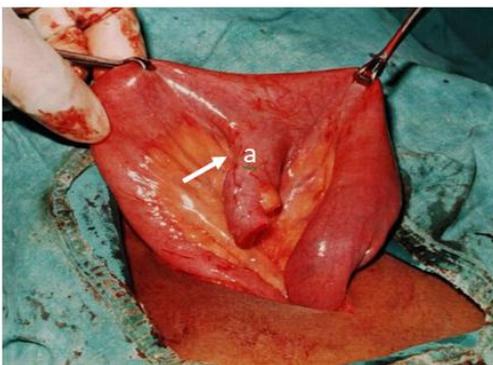


Fig. 1a. Intra operative photograph Showing a- Mesenteric location of MD of Size 4x2 cm

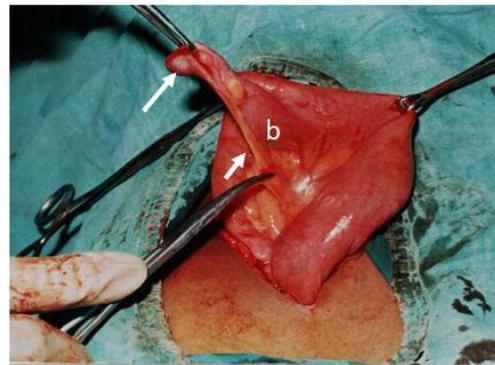


Fig. 1b. Intra operative photograph Showing a-Mesenteric MD and b-Mesodiverticular band with vitelline artery

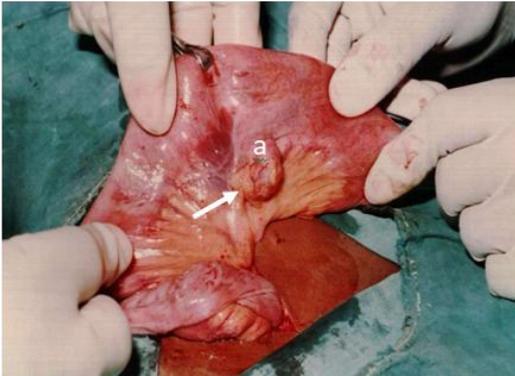


Fig. 2a. Intra operative photograph Showing a- Mesenteric location of MD of Size 2x1 cm Short and stumpy MD

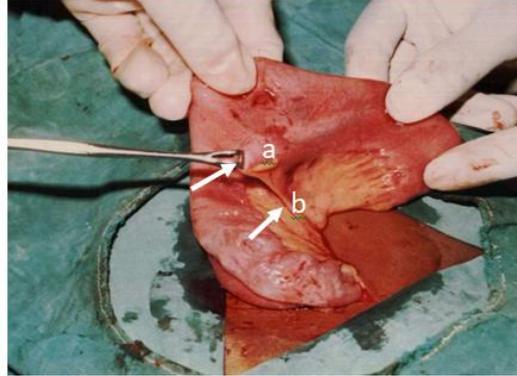


Fig. 2b. Intra operative photograph Showing a-Mesenteric MD and b-Mesodiverticular band with vitelline artery

Case II

A 27-year adult male in the year 1997, was presented in our center with complaints of pain in right McBurney's point, there was history of pain abdominal two months back. On physical examination revealed similar findings like case one. We searched perioperatively, the Meckel's Diverticulum, traced till the terminal 2 feet's of ileum proximal to ileo-caecal junction. We found a diverticulum along with the Mesenteric border of the terminal ileum, which is very rare entity.

In adult male patient having Mesenteric Meckel's Diverticulum of size 2x1 cm in length, shorts and stumpy. (Fig. 2a, 2b). In this patient there was very short vitelline artery that creates as Mesodiverticular band from the mesentery to the tip of the diverticulum, which diverts away from antimesenteric border to the root of mesentery. So it is a classical picture of Mesenteric Meckel's Diverticulum (MMD). Therefore, even though it is an asymptomatic and incidentally detected Mesenteric Meckel's Diverticulum, the surgical decision should be standard resection and anastomosis was done. The histopathology report confirmed the Meckel's diverticulum without any heterotopic tissue. Post-operative patient recovery was uneventful and patient discharge on 7th post-operative day.

3. DISCUSSION

The literature review from 1941 till now revealed 32 cases reports of mesenteric Meckel's Diverticulum. The case was distributed between 14 cases in the pediatric population and 18 cases of the adult population. The first description of a mesenteric-sided MD was

reported in 1941 by Chaffin and afterward very few cases have been reported in the surgical literature, without being documented on pre-operative imaging. In particular, Sarioglu-Buke *et al.*, offered "the possible embryological explanation that the etiology of the anomaly was due to the persistence of a short vitelline artery that creates a Mesodiverticular band from the mesentery to the tip of the diverticulum, which diverts the diverticulum away from the antimesenteric border during the elongation and growing process" [6].

Different theory's as proposed for mesenteric location of Meckel's Diverticulum are as follow.

- Persistence of short vitelline artery which creates a Mesodiverticular band from mesentery to the tip of the diverticulum, thus diverting the diverticulum away from the antimesenteric border during the process of elongation and growing process as described by Sarioglu-Buke.
- "Adherence of vitelline duct to the ileal mesentery due to congenital or inflammatory adhesions due to diverticulitis and ectopic gastric tissue" [1,2,3,5,6].

In our case, the Meckel's diverticulum was located at the mesenteric location of the ileum. "A possibility is the persistence of a very short vitelline artery that creates a Mesodiverticular band from the mesentery to the tip of the diverticulum, which diverts the diverticulum away from the antimesenteric border during rapid growth. In general, ileal duplications share the wall and the blood supply of the ileum and the Meckel's diverticulum has its own artery. However, this is still not sufficient for a differential

diagnosis because the vitelline artery is present in about 10% of cases. The anomaly presented could have been due to a short vitelline artery that disappeared without leaving a remnant or to an intrauterine adhesion between the mesentery of the ileum and the omphalomesenteric canal" [6]. Thus, during the elongation and growing process, the "stuck" diverticulum might have been diverted from the antimesenteric border of the ileum [7-10].

4. CONCLUSION

Surgeons should look for Meckel's Diverticulum not only along antimesenteric border but also seen the mesenteric border to detect the unusual location of Mesenteric Meckel's Diverticulum. Mesenteric location of Meckel's Diverticulum is more alarming because it may erode the Mesentery and its vasculature during diverticulitis causing grave complications. Incidentally detected lesions during appendectomy or laparotomy should be searched and resection anastomosis performed instead of simple wedge resection.

CONSENT

As per international standard or university standard, patient(s) written consent has been collected and preserved by the author(s).

ETHICAL APPROVAL

As per international standard or university standard written ethical approval has been collected and preserved by the author(s).

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

1. Thute PP, Bakane BC, Keche HA, et al. Mesenteric Meckel's diverticulum - a rare entity. J Evolution Med Dent Sci. 2020;9(47):3572-3574.

- DOI: 10.14260/jemds/2020/783
2. Ahmad Z, Sharma A, Vatti V, Ahmed M, Ali MA. Rare presentation on Meckels diverticulum on the mesentric border forming a mass. Int Surg J. 2014;1:188-90.
 3. Naveen Kumar C, Magesh Kumar J, Suresh Babu P, Ravishankar KS. Rare mesenteric location of Meckel's Diverticulum: A case report. International Journal of current Medical and Applied sciences. 2017;15(3):162-164.
 4. Singh A, Panda SS, Sharma N, Bajpai M. Meckel's diverticulum at uncommon mesenteric location. J Indian Assoc Pediatr Surg. 2013;18(3):127-8. DOI: 10.4103/0971-9261.116052. PMID: 24019647; PMCID: PMC3760314.
 5. Melissa M Levack, Amy G Fiedler, Haytham Kaafarani, David R King, Perforation of a mesenteric Meckel's diverticulum. Journal of Surgical Case Reports. 2018;2018(6):rjy126. Available:https://doi.org/10.1093/jscr/rjy126
 6. Sarioglu-Buke A, Corduk N, Koltuksuz U, Karabul M, Savran B, Bagci S. An uncommon variant of Meckel's diverticulum. Can J Surg. 2008;51:E46-7. [PMC free article] [PubMed] [Google Scholar]
 7. DiGiacomo JC, Cottone FJ. Surgical treatment of Meckel's diverticulum. South Med J. 1993;86:671-5.
 8. Cullen JJ, Kelly KA, Moir CR, Hodge DO, Zinsmeister AR, Melton LJ 3rd. Surgical management of Meckel's diverticulum. An epidemiologic, population-based study. Ann Surg. 1994;220(4):564-8.
 9. Vane DW, West KW, Grosfeld JL. Vitelline duct anomalies. Experience with 217 childhood cases. Arch Surg. 1987; 122:542-7. [PubMed] [Google Scholar]
 10. Maieron R, Stimac D, Avellini C, Zoratti L, Rizzi C, Scott C, et al. Acute gastrointestinal bleeding due to Meckel's diverticulum heterotopic gastric mucosa. Ital J Gastroenterol. 1996;28:225-8. [PubMed] [Google Scholar]

© 2023 Jatal et al.; This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Peer-review history:
The peer review history for this paper can be accessed here:
<https://www.sdiarticle5.com/review-history/100552>