

CASE REPORT

PEER REVIEWED | OPEN ACCESS

Barium-induced allergic appendicitis: A case report

David L. Manuel, Miriam Neufeld, Laura M. Piechura,
Qin Huang, Gentian Kristo

ABSTRACT

Appendicitis after barium administration occurs rarely and its underlying pathology remains undetermined. We present a case where acute appendicitis presented as an allergic reaction to barium sulfate. A 55-year-old male presented with nausea and severe right lower abdomen and right groin pain seven days after a barium swallow study. Computer tomography revealed retained barium in the appendix without any signs of inflammation. The patient underwent a laparoscopic appendectomy with resolution of his symptoms. Histopathologic examination demonstrated eosinophilic infiltration of the muscularis propria, consistent with acute eosinophilic appendicitis as a result of type I hypersensitivity reaction to barium. The recognition of potential risk of barium-induced allergic appendicitis is important for timely diagnosis.

Keywords: Allergic appendicitis, Barium Sulfate, Colorectal Surgery, Eosinophilic appendicitis

How to cite this article

Manuel DL, Neufeld M, Piechura LM, Huang Q, Kristo G. Barium-induced allergic appendicitis: A case report. J Case Rep Images Surg 2017;2:25–28.

Article ID: 100042Z12DM2017

doi:10.5348/Z12-2017-42-CR-7

INTRODUCTION

Barium-induced appendicitis is a rare complication after enteric barium examinations and was first reported as a case in 1954 [1]. The time span between barium study and onset of appendicitis ranges from a few hours to a few years [2] with the highest risk being within two months after barium administration [3]. Although the retained barium in the appendix is generally thought to form a barium-coated fecalith (barolith) leading to luminal obstruction and appendicitis [4], the pathophysiology of barium-induced appendicitis remains unclear. We present our experience with this rare surgical scenario in this report.

CASE REPORT

A 55-year-old male with dysphagia underwent an outpatient barium swallow study at an outside institution. A few hours later he developed nausea and severe pain in the right lower quadrant of the abdomen, prompting him to present at the same outside institution's emergency department for an evaluation. He was admitted for overnight observation after a computed tomography scan

David L. Manuel^{1,4}, Miriam Neufeld^{2,5}, Laura M. Piechura^{2,7}, Qin Huang^{3,6}, Gentian Kristo^{2,7}

Affiliations: ¹Department of Radiology, Veterans Affairs Boston Healthcare System, Boston, MA, USA; ²Department of Surgery, Veterans Affairs Boston Healthcare System, Boston, MA, USA; ³Department of Pathology, Veterans Affairs Boston Healthcare System, Boston, MA, USA; ⁴Department of Radiology, Boston Medical Center/Boston University, Boston, MA, USA; ⁵Department of Surgery, Boston Medical Center/Boston University, Boston, MA, USA; ⁶Department of Pathology, Boston Medical Center/Boston University, Boston, MA, USA; ⁷Department of Surgery, Brigham and Women's Hospital/Harvard Medical School, Boston, MA, USA.

Corresponding Author: Gentian Kristo MD, Department of Surgery, Veterans Affairs Boston Healthcare System (116C), 1400 VFW Parkway, West Roxbury, Boston, MA 02132, USA; Email: gentian.kristo@va.gov

Received: 06 March 2017

Accepted: 31 May 2017

Published: 19 June 2017

of the abdomen and pelvis which was unable to provide a diagnosis.

Six days later, the patient presented to our emergency department with nausea and severe right lower quadrant abdominal pain radiating into the right groin. Physical examination revealed significant tenderness in the right lower quadrant of the abdomen and over the right groin, without any evidence of groin hernias. The patient was afebrile, with normal leukocyte counts.

A computed tomography scan of the abdomen and pelvis was obtained and showed retained barium in the appendix without any signs of appendicitis (Figure 1).

After the patient was evaluated by consultant surgeons, he was taken immediately to the operating room for a laparoscopic exploration of the abdomen with the presumptive diagnosis of barium-induced appendicitis.

At surgery, only minimal serosal hyperemia of the body of the appendix was found and an appendectomy was performed. Postoperatively, patient reported immediate and complete relief of his right lower quadrant and right groin pain. He was discharged to home 12 hours after the surgery and had an unremarkable recovery.

Pathologic examination of the appendix showed a dilated appendiceal lumen without fecaliths, barium crystals in the mucosa of the appendix (Figure 2), absence of neutrophils in the muscle layer, and eosinophilic infiltration of the muscularis propria (Figure 3) with 150 eosinophils per high power field.

DISCUSSION

Herein, we report the case of a patient with barium-induced appendicitis, with symptoms starting hours after barium administration and correct diagnosis and treatment delayed for seven days. This indicates the importance of maintaining a high index of suspicion for

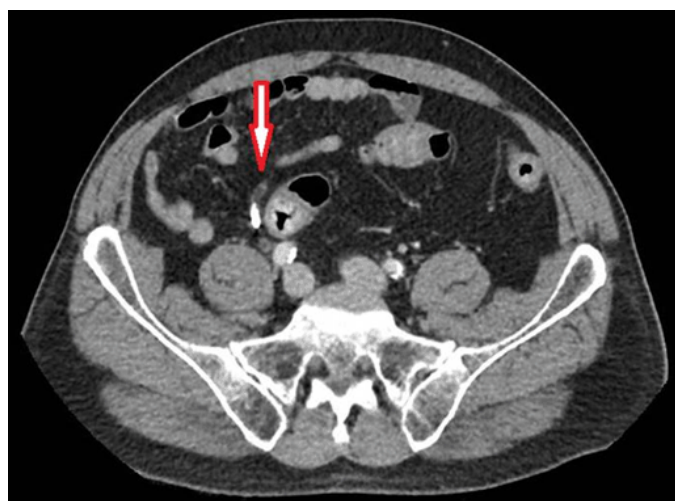


Figure 1: Axial computed tomography scan of the abdomen with IV contrast demonstrating barium retained in the appendix without edema of the appendix or inflammation in the periappendiceal fat. The tip of the appendix contains no barium and is normal (arrow).

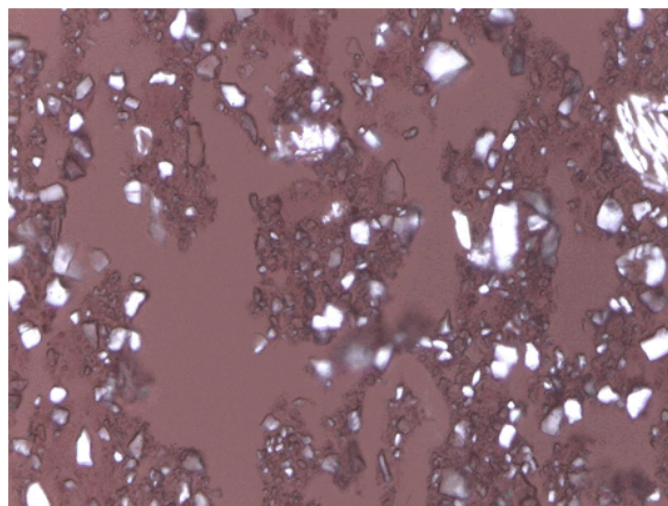


Figure 2: Polarized barium crystals deposits in the mucosa of the appendix.

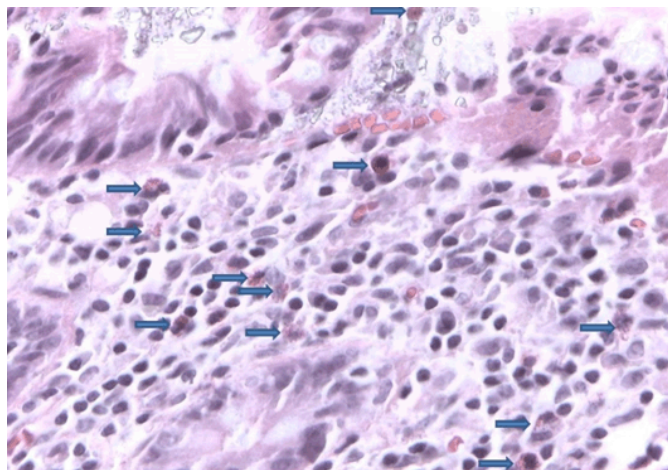


Figure 3: Eosinophilic infiltrate with degranulation (arrows) in muscularis propria.

barium-induced appendicitis in patients who present with symptoms of appendicitis after recent barium imaging.

Eosinophils are normally present in the lamina propria and submucosa, but not in the muscularis propria of the appendix [5]. The eosinophilic infiltration of the muscularis propria in our patient suggests a type I hypersensitivity reaction to barium or to any of the additives contained in the barium sulfate solution, including deflocculation agents, suspending agents, and flavoring agents [6].

If the mucosal injury caused by the eosinophils becomes infected by bacteria it leads to acute suppurative appendicitis, whereas in the absence of infection acute eosinophilic appendicitis (AEA) occurs [5].

Acute eosinophilic appendicitis is a known rare variant of appendix inflammation. The histologic hallmark of this entity is eosinophilic infiltration of the muscularis propria without neutrophilic infiltration [5, 7], as was the finding in our patient. A count of > 10 eosinophils per

high power field at microscopic examination is found in AEA [8]. In our case, there were about 150 eosinophils per high power field.

Acute eosinophilic appendicitis has been related to multiple parasites including *Strongyloides stercoralis* [9], *Schistosoma japonicum* [10], and *Entamoeba histolytica* [11].

This case report is very significant because, to the best of our knowledge, it represents the first reported evidence of an acute eosinophilic appendicitis caused by retention of the barium sulfate solution in the appendix.

CONCLUSION

Barium-induced appendicitis is a very rare clinical entity, but given the pervasive use of barium for enteric radiographic studies clinicians should recognize this potential risk to avoid delayed diagnosis and treatment. Histopathology is the gold standard for diagnosis of this rare condition.

Author Contributions

David L. Manuel – Substantial contributions to conception and design, Acquisition of data, Analysis and interpretation of data, Drafting the article, Revising it critically for important intellectual content, Final approval of the version to be published

Miriam Neufeld – Analysis and interpretation of data, Revising it critically for important intellectual content, Final approval of the version to be published

Laura M. Piechura – Analysis and interpretation of data, Revising it critically for important intellectual content, Final approval of the version to be published

Qin Huang – Analysis and interpretation of data, Revising it critically for important intellectual content, Final approval of the version to be published

Gentian Kristo – Analysis and interpretation of data, Revising it critically for important intellectual content, Final approval of the version to be published

Guarantor

The corresponding author is the guarantor of submission.

Conflict of Interest

Authors declare no conflict of interest.

Copyright

© 2017 David L. Manuel et al. This article is distributed under the terms of Creative Commons Attribution License which permits unrestricted use, distribution and reproduction in any medium provided the original author(s) and original publisher are properly credited. Please see the copyright policy on the journal website for more information.

REFERENCES

1. Gubler JA, Kukral AJ. Barium appendicitis. J Int Coll Surg 1954 Mar;21(3 1):379–84.
2. Novotny NM, Lillemoe KD, Falimirski ME. Barium appendicitis after upper gastrointestinal imaging. J Emerg Med 2010 Feb;38(2):148–9.
3. Li HM, Yeh LR, Huang YK, Lin CL, Kao CH. The association between barium examination and subsequent appendicitis: A nationwide population-based study. Am J Med 2017 Jan;130(1):54–60.e5.
4. Maglinte DD, Bush ML, Aruta EV, Bullington GE. Retained barium in the appendix: Diagnostic and clinical significance. AJR Am J Roentgenol 1981 Sep;137(3):529–33.
5. Aravindan KP, Vijayaraghavan D, Manipadam MT. Acute eosinophilic appendicitis and the significance of eosinophil - Edema lesion. Indian J Pathol Microbiol 2010 Apr–Jun;53(2):258–61.
6. O'Connor SD, Summers RM. Revisiting oral barium sulfate contrast agents. Acad Radiol 2007 Jan;14(1):72–80.
7. Aravindan KP. Eosinophils in acute appendicitis: Possible significance. Indian J Pathol Microbiol 1997 Oct;40(4):491–8.
8. Carr NJ. The pathology of acute appendicitis. Ann Diagn Pathol 2000 Feb;4(1):46–58.
9. Noodleman JS. Eosinophilic appendicitis: Demonstration of *Strongyloides stercoralis* as a causative agent. Arch Pathol Lab Med 1981 Mar;105(3):148–9.
10. Kanoksil W, Larbcharoensub N, Soontrapa P, Phongkitkarun S, Sriphojanart S, Nitiyanant P. Eosinophilic appendicitis caused by *Schistosoma japonicum*: A case report and review of the literature. Southeast Asian J Trop Med Public Health 2010 Sep;41(5):1065–70.
11. Egeli T, Okudan M, Taskesen F, Celik SY, Tasdemir N. Acute eosinophilic appendicitis: An unusual variant of appendix inflammation. Turk J Colorectal Dis 2013;23(2):107–10.

Access full text article on
other devices



Access PDF of article on
other devices

