

Myxoglobulosis in the appendix

Apendiks miksglobulozisi

Cengiz Koçak¹, Akile Zengin², İbrahim Girgin³, Fatma Ferda Kartufan⁴, Mehmet Hüseyin Metineren¹

ABSTRACT

Myxoglobulosis is a specific type of mucocoele consisting of mucoid material. It is characterized by opaque, transparent globules that resemble "fish eggs" or "frogspawns". It is generally diagnosed incidentally during an appendectomy or an autopsy. In this paper, we aim to present the case of a 58-year-old male patient who was referred to our hospital because of abdominal pain and loss of appetite. The patient underwent an appendectomy. Opaque intraluminal globules were found in the appendectomy material. The globules resembled pearls and they were 2–3 mm in diameter. After histopathological examinations, the patient was diagnosed with myxoglobulosis accompanied with acute appendicitis. According to our research, this is the first case of myxoglobulosis in our country.

Keywords: Acute appendicitis, myxoglobulosis, appendectomy

ÖZET

Miksoglobulozis mukoid materyal içeren spesifik bir mukosel tipidir. Balık yumurtası veya kurbağa yumurtasına benzeyen opak, şeffaf globüller ile karakterizedir. Genellikle apendektomi veya otopsi sırasında tesadüfi olarak teşhis edilmektedir. Bu yazımızda hastanemize karın ağrısı ve iştahsızlık ile başvuran ve apendektomi yapılan 58 yaşında erkek hastada saptanmış apendiks miksglobulozisi vakasını sunuyoruz. Apendektomi materyali içinde opak, 2-3 mm çapında, inciye benzeyen intraluminal globüller izlendi. Histopatolojik inceleme sonrasında hastaya akut apandisit eşlik eden miksglobulozis tanısı konuldu. Bizim araştırmalarımıza göre bu vaka bizim ülkemizden bildirilen ilk miksglobulozis vakasıdır.

Anahtar Kelimeler: Akut apandisit, miksglobulozis, apendektomi

INTRODUCTION

Myxoglobulosis or caviar appendix is a special type of mucocoele. It is characterized by the presence of opaque globules. Globules are made up of mucoid material and they form clusters that resemble "fish eggs" or "frogspawns." The first case of myxoglobulosis was described by Latham in 1897, who found the condition in a postmortem examination, as reported in the manuscript of Probst and Lassar (1). In 1914, the term "myxoglobulosis" was used by von Hansemann, who analyzed opaque globules varying in diameter from 0.1 to 1.0 cm (2), and it was described as a variant of mucocoele. The etiology and pathogenesis of myxoglobulosis are still uncertain. Probst and Lassar (1) have reported that the probable etiological factors are bacteria and necrotic epithelial debris, which may cause nidus formation for the deposition of mucin. Lubin and Berle (2) have reported that the core of the globules represent a mass of mucin by the granulation tissue that originated from the appendiceal wall and then these globules are disintegrated by mechanical contractions. Li et al. (3) have reported that the globules occur as a formation of mucin by the granulation tissue capsule and then these globules are extruded into the lumen by mechanical forces.

In this case report, we present an extremely rare case of appendiceal myxoglobulosis. According to our studies, this case is the first report of myxoglobulosis in our country.

CASE PRESENTATION

The procedures of this case were performed in accordance with the Declaration of Helsinki. The patient was instructed and then informed consent was received. A 58-year-old male patient was referred to the Gediz State Hospital in July 2012 with complaints of nausea and abdominal pain in the right lower abdomen. His historical and anamnestic data were not unusual, but he had a history of smoking for 46 years and had been diagnosed with chronic obstructive pulmonary disease. Physical examination revealed that he had rebound tenderness. His blood cell count tests showed that the number of white blood cells had slightly increased. Based on these findings, the patient was diagnosed with acute appendicitis. After spinal anesthesia was performed, the patient underwent an appendectomy. During the operation, the appendix was enlarged and distended. Then, the appendix was removed and appendectomy material was sent to our pathology laboratory for histopathological examination.

Macroscopic examination results revealed that the appendectomy specimen was 7 cm in length and 1 cm in thickness. In the area that was cut, the appendiceal lumen was dilated and filled with numer-

¹Department of Pathology, Dumlupınar University Faculty of Medicine, Kütahya, Turkey

²Department of General Surgery, Dumlupınar University Faculty of Medicine, Kütahya, Turkey

³Clinic of General Surgery, Gediz State Hospital, Kütahya, Turkey

⁴Clinic of Anesthesiology, Gediz State Hospital, Kütahya, Turkey

Address for Correspondence Yazışma Adresi

Akile Zengin

Dumlupınar Üniversitesi Tıp Fakültesi, Genel Cerrahi Anabilim Dalı, Kütahya, Türkiye
Phone: +90 274 265 20 31
e-mail: dr.akile.zengin@gmail.com

Received / Geliş Tarihi: 23.12.2014
Accepted / Kabul Tarihi: 29.03.2015
Available Online Date / Çevrimiçi Yayın Tarihi: 14.07.2015

©Copyright 2015
by Turkish Surgical Association
Available online at
www.ulusalcerahidergisi.org

©Telif Hakkı 2015
Türk Cerrahi Derneği

Makale metnine
www.ulusalcerahidergisi.org
web sayfasından ulaşılabilir.



Figure 1. Numerous pearl-like globules in the dilated appendix lumen



Figure 2. Macroscopic appearance of the globules resembling "fish eggs"

ous opaque, whitish pearl-like globules 2–3 mm in diameter (Figure 1). The accumulation of these globules resembled fish eggs (Figure 2). Biopsy samples were examined histopathologically. On microscopic examination of samples stained by hematoxylin and eosin (H&E), the globules which consisted of eosinophilic laminations of mucin surrounding an amorphous granular core were observed. The diffusion of inflammatory infiltration consisting of polymorphonuclear leucocytes and mononuclear cells were observed in the entire layer of the appendiceal wall. Furthermore, hemorrhage was evidently observed in the wall. Globules were positively stained with Alcian Blue (Figure 3, 4). Considering all these histopathological findings, our case was diagnosed as myxoglobulosis accompanied with acute appendicitis.

DISCUSSION

The incidence of the appendiceal mucocoele is estimated to be 0.2%–0.3% of the appendectomy specimens, with myxoglobulosis constituting 0.35%–0.8% of mucocoeles (4–6). There are patients with myxoglobulosis who clinically exist in a manner similar to those with a mucocoele of the appendix. The disorder occurs most commonly in the sixth or seventh decade of life. Patients are generally asymptomatic, and myxoglobulosis is found by chance during autopsies or laparotomies for other reasons. Sometimes, right lower abdominal pain and acute appendicitis is observed in patients with myxoglobulosis (7). To our knowledge, this case is the first report of myxoglobulosis in our country. The development of myxoglobulosis requires

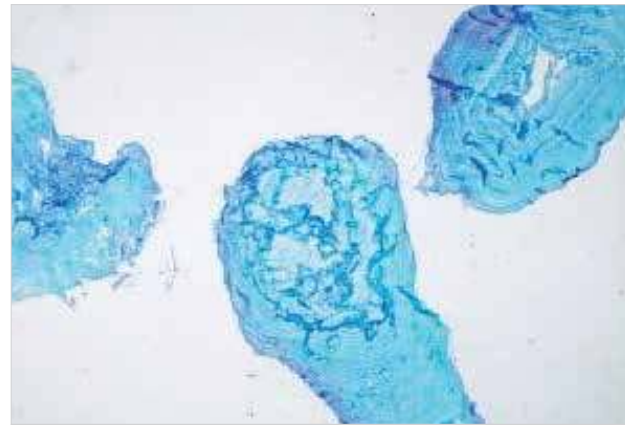


Figure 3. Alcian Blue positive staining of the globules (Alcian Blue pH 2.5×20)

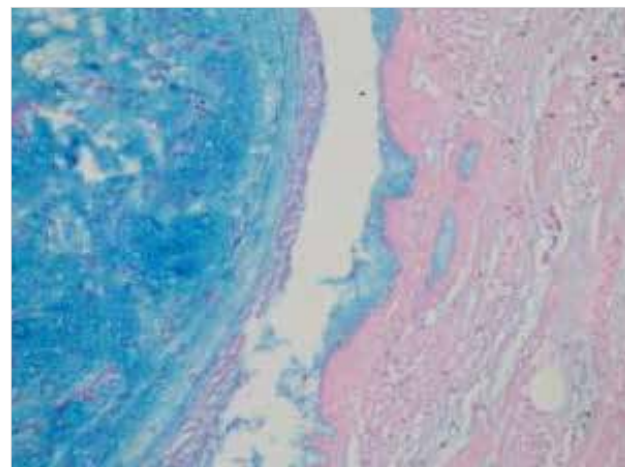


Figure 4. Alcian Blue positive staining of the globules and adjacent appendiceal luminal mucosa (Alcian Blue pH 2.5×40)

obstruction to the proximal appendiceal lumen with continued production of mucin distally. The causes of the proximal appendiceal obstruction include fecalith, epithelial hyperplasia, post-inflammatory fibrosis, cystadenoma, cystadenocarcinoma, carcinoid, and endometriosis. The characteristic feature is the presence of opaque, pearl-like globules in the appendiceal lumen. The most frequent complication is peritonitis or pseudomyxoma peritonei (8, 9). In the literature, the number of reported cases of myxoglobulosis is extremely rare. Recently, Aroukatos et al. (10) reported that a case of myxoglobulosis of the appendix is associated with a ruptured diverticulum. Falah et al. (11) have reported a case of appendiceal myxoglobulosis associated with peritonitis due to perforated peptic ulcer. Padhy et al. (12) have reported a case of myxoglobulosis of the appendix. Routine histopathological examination is essential to diagnose myxoglobulosis.

CONCLUSION

In this paper, we have reported a rare case of myxoglobulosis, which was incidentally found in a patient with peritonitis due to acute appendicitis.

Informed Consent: Written informed consent was obtained from patient who participated in this case.

Peer-review: Externally peer-reviewed.

Author Contributions: Concept - C.K., I.G., F.F.K.; Design - A.Z., M.H.M.; Supervision - I.G.; Funding - A.Z.; Materials - M.H.M.; Data Collection and/or Processing - I.G.; Analysis and/or Interpretation - M.H.M.; Literature Review - C.K., A.Z., F.F.K.; Writer - A.Z.; Critical Review - C.K., F.F.K.

Conflict of Interest: No conflict of interest was declared by the authors.

Financial Disclosure: The authors declared that this study has received no financial support.

Hasta Onamı: Yazılı hasta onamı bu olguya katılan hastadan alınmıştır.

Hakem Değerlendirmesi: Dış bağımsız.

Yazar Katkıları: Fikir - C.K., I.G., F.F.K.; Tasarım - A.Z., M.H.M.; Denetleme - I.G.; Kaynaklar - A.Z.; Malzemeler - M.H.M.; Veri toplanması ve/veya işlemesi - I.G.; Analiz ve/veya yorum - M.H.M.; Literatür taraması - C.K., A.Z., F.F.K.; Yazıyı yazan - A.Z.; Eleştirel inceleme - C.K., F.F.K.

Çıkar Çatışması: Yazarlar çıkar çatışması bildirmemişlerdir.

Finansal Destek: Yazarlar bu çalışma için finansal destek almadıklarını beyan etmişlerdir.

REFERENCES

1. Probststein JG, Lassar GN. Mucocele of the appendix, with myxoglobulosis. *Ann Surg* 1948; 127: 171-176. [\[CrossRef\]](#)
2. Lubin J, Berle E. Myxoglobulosis of the appendix. Report of two cases. *Arch Pathol* 1972; 94: 533-536.
3. Li TJ, Kitano M, Yoshida A, Iwashige Y, Yamashita S. Myxoglobulosis is an extravasation mucocele of the lower lip. *J Oral Pathol Med* 1997; 26: 342-344. [\[CrossRef\]](#)
4. Gonzalez JE, Hann SE, Trujillo YP. Myxoglobulosis of the appendix. *Am J Surg Pathol* 1988; 12: 962-966. [\[CrossRef\]](#)
5. Rajiman I, Leong S, Hassaram S, Marcon NE. Appendiceal mucocele: endoscopic appearance. *Endoscopy* 1994; 26: 326-328. [\[CrossRef\]](#)
6. Madwed D, Mindelzun R, Jeffrey RB Jr. Mucocele of the appendix: imaging findings. *AJR Am J Roentgenol* 1992; 159: 69-72. [\[CrossRef\]](#)
7. Viswanath YK, Griffiths CD, Shipsey D, Oriolowo A, Johnson SJ. Myxoglobulosis, a rare variant of the appendiceal mucocele, occurring secondary to an occlusive membrane. *J R Coll Surg Edinb* 1998; 43: 204-206.
8. Brustmann H. Myxoglobulosis of the appendix associated with a proximal carcinoid and a pseudodiverticulum. *Ann Diagn Pathol* 2006; 10: 166-168. [\[CrossRef\]](#)
9. Alcalay J, Alcalay L, Lorent T. Myxoglobulosis of the appendix. *Br J Radiol* 1985; 58: 183-184. [\[CrossRef\]](#)
10. Aroukatos P, Verras D, Vondoros GP, Repanti M. Myxoglobulosis of the appendix: a case associated with ruptured diverticulum. *Case Rep Med* 2010; 2010: 745021. [\[CrossRef\]](#)
11. Falah HH, Maafi AA, Chatrnour G, Jahromi SK, Ebrahimi H. Myxoglobulosis, a rare variant of appendiceal mucocele : case report and review of the literature. *Thrita J Med Sci* 2013; 2: 155-158.
12. Padhy BP, Panda SK. Myxoglobulosis of appendix a rare entity. *Indian J Surg* 2013; 75: 337-339. [\[CrossRef\]](#)