

# AMSER Case of the Month

## December 2023

HPI 60 year old female presenting with lower abdominal  
pain

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# Patient Presentation

**HPI:** 60 year old female presenting with lower abdominal pain.

**PMH:** Hypertension, Irritable Bowel Syndrome, Osteopenia, Pancreatitis, and Ulcerative Colitis.

**Medications:** amlodipine and sulfasalazine

**Vitals:** BP: 138/80, Pulse: 70, Resp: 16, SpO2: 98%

**Physical Exam:**

- Genitourinary: No CVA tenderness
- Gastrointestinal: Soft, lower abdominal tenderness, non-distended, and positive bowel sounds.

# Patient Presentation

## Labs:

### -CBC:

White Blood Cell	Red Blood Cell	Hemoglobin	MCV	Platelets
6.2 (normal)	4.53 (normal)	14.0 (normal)	92 (normal)	209 (normal)

### -BMP:

Glucose	BUN	Creatinine	Sodium	Potassium	Chloride	CO2	Calcium	GFR
108 (normal)	8 (normal)	0.63 (normal)	141 (normal)	4.3 (normal)	103 (normal)	26 (normal)	9.4 (normal)	102 (normal)

What Imaging Should We Order?

# Select the applicable ACR Appropriateness Criteria

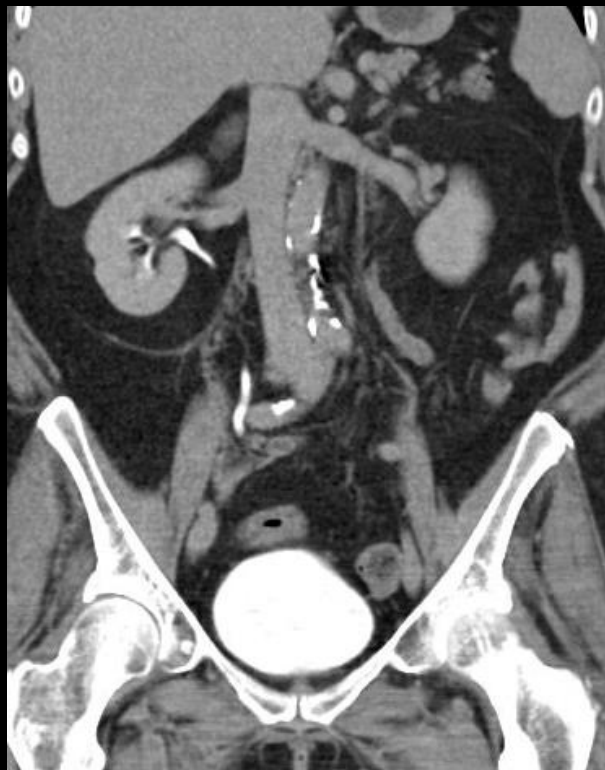
**Variant 1:** Right lower quadrant pain. Initial imaging.

Procedure	Appropriateness Category	Relative Radiation Level
CT abdomen and pelvis with IV contrast	Usually Appropriate	⊕⊕⊕
US abdomen	May Be Appropriate	○
US pelvis	May Be Appropriate	○
MRI abdomen and pelvis without and with IV contrast	May Be Appropriate	○
MRI abdomen and pelvis without IV contrast	May Be Appropriate	○
CT abdomen and pelvis without IV contrast	May Be Appropriate	⊕⊕⊕
Radiography abdomen	Usually Not Appropriate	⊕⊕
Fluoroscopy contrast enema	Usually Not Appropriate	⊕⊕⊕
CT abdomen and pelvis without and with IV contrast	Usually Not Appropriate	⊕⊕⊕⊕
WBC scan abdomen and pelvis	Usually Not Appropriate	⊕⊕⊕⊕

This imaging modality was ordered by the primary care physician



# Findings: (unlabeled)





# Findings: (labeled)

blind ending pouch originating at the cecum

Punctate and curvilinear wall calcifications within a tubular low attenuating mass continuous with the cecal base.

Sagittal

Coronal

Coronal

Coronal

Cecum

Appendix

**Final Dx:**

Appendiceal Mucocele



# Case Discussion: appendiceal mucocele

*Appendiceal mucocele is distention of the appendix due to mucous accumulation in the lumen. It is a general term used to describe its appearance on radiological findings, and has many possible causes [1].*

**Presentation:** Patient may experience abdominal pain, tenderness to palpation, nausea, vomiting, and/or fever [2, 3]. A quarter of patients are asymptomatic. Hematuria and urinary dysfunction, quite rare, can also be seen [4].

**Imaging:** On radiograph, appears as mass in right iliac fossa with peripheral calcifications. CT shows spherical mass with well-defined edges that shares border with cecum [1].

**Pathology:** Common causes include mucus retention cyst from obstruction of mucous sac, serrated polyp, and mucinous neoplasms [1]. There are three main types of histological groups consisting of mucinous cystadenocarcinoma, mucinous cystadenoma, and nonneoplastic mucocele.

# Case Discussion: appendiceal mucocele

**Pathology cont:** Mucinous cystadenocarcinoma has glandular and stromal invasion and is similar to mucinous colonic tumors [5].

**Epidemiology:** Appendiceal mucocele is a rare condition found only 0.2-0.3% of the time at appendectomy [1, 2, 3]. It appears in middle-aged populations the most [1]. Patients are six times more likely to develop an adenocarcinoma [7].

**Management:** Treated with surgery; a hemicolectomy is done if malignancy is suspected [1, 6]. Surgery has good long-term prognosis [6].

**Complications:** Can cause internal bleeding, obstruction, melena, or pyonephrosis [6]. Pseudo-myxoma peritonei syndrome occurs when there is extra appendicular spread into the peritoneal cavity. It consists of gelatinous material composed of a combination of mucin, mucinous epithelium, and adenocarcinoma.

# Case Discussion: appendiceal mucocele

**Complications cont:** The composition of the gelatinous material will be ultimately diagnosed as peritoneal adenomucinosis (benign) versus peritoneal mucinous carcinomatosis, which originates from appendiceal mucinous adenocarcinoma. There is an intermediate category which contains both histologies. The 5 year survival rate for patients with pseudo-myxoma peritonei syndrome is 84% for adenomucinosis, 37.5% for the intermediate category, and 6.7% for mucinous carcinomatosis [8].

# References:

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