Spontaneous Recurrent Pneumoperitoneum in Systemic Sclerosis

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Key Words
Recurrent pneumoperitoneum · Systemic sclerosis · Abdominal distention · Constipation

Case Report
A 72-year-old male presented with intermittent painful abdominal distension and constipation. He explained that the symptoms occurred during the past 2 years, with variable intervals (>4 episodes per year) characterized by diffuse abdominal pain, abdominal distention, vomiting, and diarrhea, followed by constipation; these episodes had variable duration, between 5 and 15 days, and resolved spontaneously. The patient denied dysphagia, fecaloid vomiting, weight loss, or hemorrhage. Recent upper gastrointestinal endoscopy and ileocolonoscopy were normal. His past medical history included systemic sclerosis diagnosed at 56 years of age and treated with methotrexate for the past 7 years with improvement of cutaneous and articular symptoms and an exploratory laparotomy when he was 71 years old due to pneumoperitoneum with no identifiable cause. Other regular medications, smoking, or alcohol consumption were also denied. Family history was irrelevant. On admission, the patient was stable and without fever; the abdomen was markedly distended, with decreased bowel sounds, diffuse tympanism, and tenderness but without peritoneal signs. Blood tests showed no significant abnormalities. Standing chest and abdominal radiographs revealed free air under both hemidiaphragms, distension of the small and large intestine, and air-fluid levels (Fig. 1, 2). Abdominal and pelvic contrast-enhanced computed tomography confirmed the presence of pneumoperitoneum and showed many distended loops of the small bowel, but no cause of the free air was identified (Fig. 3, 4).

The patient’s general condition was stable, and antibiotics, analgesics, and promotility agents were started. Despite the absence of other causes and due to the chronic presentation, pneumoperitoneum was considered a manifestation of systemic sclerosis. The patient was discharged on day 7 with symptomatic medication (butylscopolamine, domperidone, and simethicone).

Six months later, abdominopelvic computed tomography was repeated and revealed a smaller pneumoperitoneum and the presence of small air bubbles in the colonic wall; no obstructive lesion was identified. The symptoms improved with oral medication.

One year later, magnetic resonance enterocolonography showed mild pneumoperitoneum, normal contrast progression, and distension of the small intestine; there were no stenotic lesions, fistulas, or contrast extravasation into the pneumoperitoneum. At outpatient follow-up 24 months later, the patient had recurrent mild abdominal distension but remained clinically well. In this case, the most suitable cause of the nonsurgical recurrent pneumoperitoneum was systemic sclerosis.
Fig. 1. Standing abdominal radiograph showing pneumoperitoneum, intestinal distension, and air-fluid levels.

Fig. 2. Chest radiograph revealing free air under both hemidiaphragms (pneumoperitoneum).

Fig. 3. Abdominal contrast-enhanced computed tomography showing pneumoperitoneum (sagittal view).

Fig. 4. Abdominal contrast-enhanced computed tomography showing pneumoperitoneum and many distended loops of the small bowel (axial view).
**Discussion**

Pneumoperitoneum is often caused by visceral perforation, with symptoms of peritonitis, and requires an emergent surgical intervention [1]. Nonsurgical spontaneous pneumoperitoneum is a rare entity not associated with organ perforation and usually treated conservatively [1, 2].

Nonsurgical pneumoperitoneum is associated with postoperatively retained air, ventilatory support, spontaneous rupture of pulmonary blebs, cardiopulmonary resuscitation, endoscopy, diverticulosis, pneumatosis cystoides intestinalis, pneumocholecystitis, collagen vascular diseases, gynecologic causes, iatrogenic, and idiopathic causes [2]. In our case, systemic sclerosis was the apparent etiology [1, 3]. There are few reports of chronic scleroderma-related pneumoperitoneum [1–3].

The excessive fibrosis, inflammation, and vascular dysfunction characteristic of systemic sclerosis can affect any site within the gastrointestinal tract [1]. In the small intestine, it causes hypomotility, pseudo-obstructions, and luminal distension [1]. Nevertheless, the pathogenesis of this condition is uncertain, and treatment is usually supportive (promotility and probiotic agents, laxatives, and rifaximin) [1].

Most patients with nonsurgical pneumoperitoneum have clinical clues or associated conditions that may indicate the possibility of this diagnosis. When signs of peritoneal irritation, fever, and leukocytosis are absent, nonsurgical causes of pneumoperitoneum should be considered, and conservative management with supportive care and careful observation is indicated [1–3]. Thus, recognized, nonsurgical pneumoperitoneum allows avoidance of an unnecessary laparotomy and its associated morbidity [1–3].

**Statement of Ethics**

This study did not require informed consent or review/approval by the appropriate ethics committee.

**Disclosure Statement**

The authors have no conflicts of interest to declare.

**References**