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Pneumatosis Intestinalis: A Case Report



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**Meher Angez Rahman*¹, Khaled A. H. Fahid¹, Hjh
Norrul Ainun Hj Md Thani¹, Mizanul Hasan²**

¹*Department of Radiology and Imaging, Suri Seri
Begawan Hospital, Kuala belait, Brunei.*

²*Department of Ultrasound, Popular Diagnostic Centre
Ltd., Dhaka, Bangladesh.*

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ABSTRACT

Pneumatosis intestinalis is a condition in which multiple gas-filled cysts are located in the bowel wall. It can represent a wide spectrum of disease and a variety of underlying diagnoses. The present report describes the case of a 58-year-old man with the complaints of abdominal distension, fever, vomiting and abdominal pain on and off. He had a history of moderately differentiated adenocarcinoma of rectum, Stage IIC (T3N2b) and anterior resection done.



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INTRODUCTION

Pneumatosis intestinalis (PI) is a finding characterized by the presence of gas within the bowel wall (1,2) small or large intestine. Intramural gas can affect the stomach known as gastric pneumatosis(3). Pneumatosis intestinalis has appeared in the literature as pneumatosis cystoides intestinalis, intramural gas, pneumatosis coli, pseudo lipomatosis, intestinal emphysema, bullous emphysema of the intestine and lymphopneumatosis (4,5). Pneumatosis intestinalis usually found with the condition that disrupt mucosal integrity, such as necrotizing enterocolitis, intestinal ischaemia, inflammatory bowel disease, and intestinal infections.

The incident of pneumatosis intestinalis is difficult to ascertain since most patients are asymptomatic and never come to clinical attention (5). Adults are typically diagnosed in the fifth to eighth decade.

Case report

The patient, a 58 years old man, presented with abdominal pain and distension at outpatient department of Suri Seri Begawan Hospital at Kuala belait, Brunei. On examination abdomen was mildly distended and tender, soft, bowel sound was present. Plain X-ray abdomen showed no free gas, small bowel loops were moderately dilated. Blood tests showed increased WBC, CRP and prothrombin time, de arranged liver function tests. Contrast enhanced CT scan of abdomen and pelvis revealed no free gas, dilated small bowel (jejunum), intramural gas in small bowel, ascending and descending colon, mild to moderate ascites, thrombus in portal, superior mesenteric and inferior mesenteric veins and their proximal branches. Portal (21mm) and superior mesenteric (20mm) veins were mildly dilated. Upper GI tract endoscopy showed diffuse gastritis and oesophageal candidiasis. He had a history of moderately differentiated adenocarcinoma of rectum, Stage IIC (T3N2b) and anterior resection was done. However, no recurrence of lesion was seen.

Patient was treated conservatively and discharged.

Regular SOPD (surgical outpatient department) follow-up showed no complain and no recurrence of abdominal pain.



Fig-A

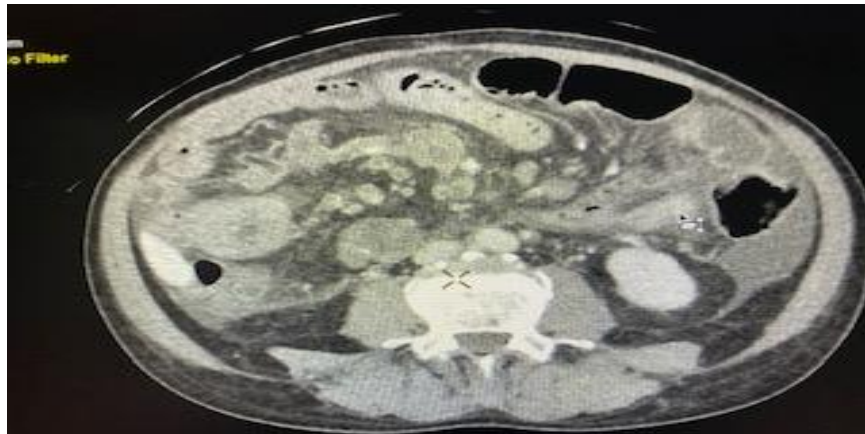


Fig-B

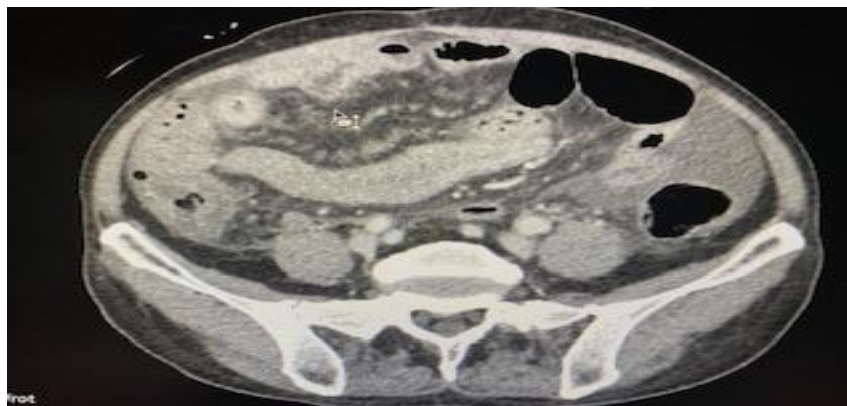


Fig C

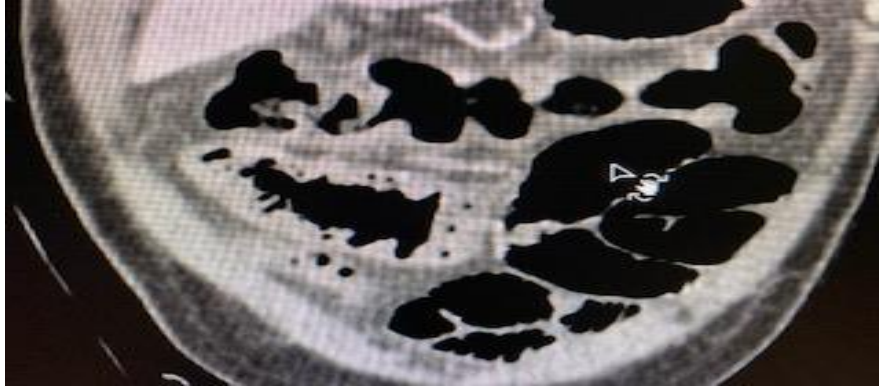


Fig-D



HUMAN

Fig-E

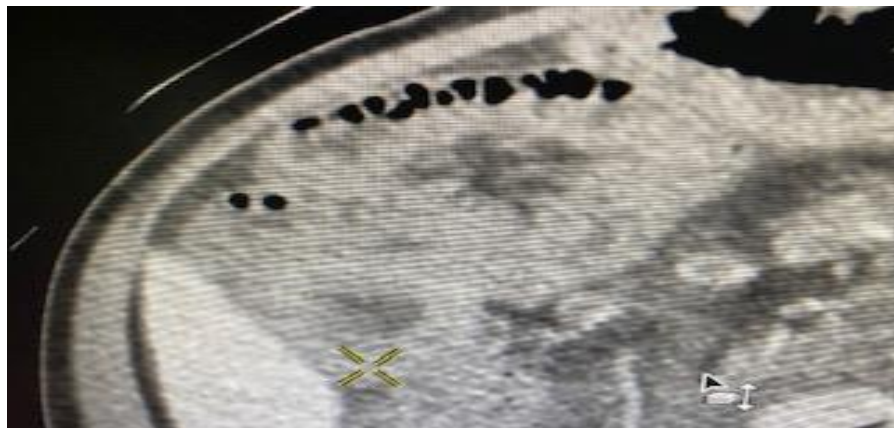


Fig-F

Fig-A to F, CECT scan of abdomen showing pneumatosis intestinalis.

DISCUSSION

Pneumatosis intestinalis (PI) is idiopathic (15 percent) or secondary (85 percent) to a variety of gastrointestinal and non-gastrointestinal illness (6,7).

A mechanical theory hypothesizes that gas dissects in to the bowel wall from either the intestinal lumen (breaks in the mucosa or through serosal surface by tracking along mesenteric blood vessels) (8). From bowel wall gas may spread along the mesentery to distant sites (9). Or from the lungs (obstructive pulmonary disease) via rupture of alveoli due to coughing air track along blood vessels in to the mediastinum, through the diaphragm, and ultimately to the mesenteric root to mesenteric blood vessels and penetrate to bowel wall.

A bacterial theory proposes that gas-forming bacilli enter the submucosa through mucosal rents or increased mucosal permeability and produce gas within the bowel wall (9,10).

Biochemical theory proposes that luminal bacteria produce excessive amount of hydrogen gas produces pressure within the intestinal lumen, gas may force directly through the mucosa and become trapped within the submucosa (11).

PI has been reported in patients with small bowel bacterial overgrowth and in patients taking alpha glucosides inhibitors which increases intestinal gas (12-16).

Most patients with pneumatosis intestinalis (PI) are asymptomatic and probably never come to clinical attention (5). Patients who come with clinical attention presents with abdominal pain, distension, vomiting, diarrhoea, weight loss and loss of appetite.

Imaging play an important role in diagnosis. In abdominal X-ray intramural gas can appear as linear, curvilinear or circular appearances. Pneumoperitonium or dilated bowel loops can be seen. An abdominal ultrasound may reveal bowel wall gas echoes with acoustic shadowing. Contrast enhanced CT scan can confirm the diagnosis and underlying causes. Usually intramural gas, free gas in peritoneum, portal venous gas, dilated bowel loops are the common findings. (17).

In upper GI tract endoscopy submucosal cysts may appear as pale or bluish in appearance.

Laboratory studies usually show leucocytosis, elevated haematocrit, metabolic acidosis and increased serum lactate.

Complication is less likely and are intestinal obstruction, volvulus, intussusception, adhesion, hematochezia and pneumoperitonium.

CONCLUSION

Exploratory laparotomy is done. If there are signs of peritonitis, metabolic acidosis, lactate >2.0mmol/L, portal venous gas. However, our case was successfully managed conservatively.

Localization of lesions and identifications of PI patterns; integration of imaging, laboratory and clinical findings permit clinicians to suspect the onset of complications and to distinguish benign from life threatening PI and to decide proper management.

Conflict of interest: None.

REFERENCES

1. G.H.Micklefield,H.D.Kuntz, and B.May, "Pneumatosis cystoides intestinalis: case reports and review of the literature",Materia Medica Polona,vol.22,no.2.pp.70-72,1990.
2. S.D. St. Peter, M.A. Abbas, and K.A. Kelly, "The spectrum of pneumatosis intestinalis", Archives of surgery, vol.138, no.1, pp.68-75,2003.
3. Cordum NR, Dixon A, Campbell DR. Gastroduodenal pneumatosis: endoscopic and histological findings. Am J Gastroenterol 1997;92:692.
- 4.Sachse RE, Burkey GW 3rd,Jonas M, et al. Benign pneumatosis intestinalis with subcutaneous emphysema in a liver transplant recipient. Am J Gastroenterol 1990;85:876.
5. Heng Y,Schuffler MD,Hanggit RC,Rohrmann CA. pneumatosis intestinalis:a review. Am J Gastroenterol 1995;90:1747.
6. Koss LG. Abdominal gas cysts (pneumatosis cystoidesintestinatorum hominis); an analysis with a report of a case and a critical review of the literature. AMA Arch Pathol 1952;53:523.
7. Knechtle SJ,Davidoff AM, Rice RP. Pneumatosis intestinalis. Surgical management and clinical outcome. Ann Surg 1990; 212:160.
8. Pieterse AS,Leong AS, Rowland R. The mucosal changes and pathogenesis of pneumatosis cystoidesintestinalis.HumPathol 1985;16:683.
9. B.L. Pear, "Pneumatosis intestinalis: a review", Radiology, vol.207, no.1, pp.13-20,1998.
- 10.S. Galandiuk and V.W. Fazio, "Pneumatosis cystoides intestinalis a review of the literature" Disease of the colon and rectum, vol.29, no.5, pp.358-363,1986.
11. Sartor RB,Murphy ME,Rydzak E. Miscellaneous inflammatory and structural disorders of the colon. In: Textbook of Gastroenterology, 3rd ed, Yamada T, Alpers D, Laine L, et al (Eds), Lippincott Williams & Wilkins, Philadelphia 1999.Vol 1877.
12. Levitt MD, Olsson S. Pneumatosis cystoides intestinalisand high breath H2 excretion: insights into the role of H2 in this condition. Gastroenterology 1995;108:1560.

13. Hisamoto A, Mizushima T, Sato K, et al. Pneumatosis cystoides intestinalis after alpha-glucosidase inhibitor treatment in a patient with intestinal pneumonitis. *Intern Med* 2006;45:73.
14. Hayakawa T, Yoneshima M, Abe T, Nomura G. Pneumatosis cystoides intestinalis after treatment with an alpha-glucosidase inhibitor. *Diabetes Care* 1999; 22:366.
15. Yanaru R, Hizawa K, Nakamura S, et al. Regression of pneumatosis cystoides intestinalis after discontinuing of alpha-glucosidase inhibitor administration. *J Clin Gastroenterol* 2002;35:204.
16. Azami Y. Paralytic ileus accompanied by the pneumatosis cystoides intestinalis after acarbose treatment in an elderly diabetic patient with a history of heavy intake of maltitol. *Intern Med* 200;39:826.
17. Olson DE, Kim YW, Ying J, Donnelly LF, CT predictors for differentiating benign and clinically worrisome pneumatosis intestinalis in children beyond the neonatal period. *Radiology* 2009; 253:513.

