

Ascites Caused by Intestinal Anisakiasis: A Case Report and Literature Review

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Anisakiasis is a parasitic infection affecting the human gastrointestinal tract. It is caused by the consumption of contaminated, raw or inadequately cooked fish or squid, which is typically used for making sushi and sashimi. Most cases involve gastric anisakiasis, whereas intestinal anisakiasis is rare. This report describes the case of a 63-year-old Japanese woman with a history of raw fish consumption who presented with acute-onset abdominal pain and vomiting. Abdominal computed tomography (CT) demonstrated thickened small bowel loops and ascites on the liver surface. The patient was admitted for supportive care. On the second day of hospitalization, contrast-enhanced abdominal CT revealed that the ascites had moved from the liver surface to the pouch of Douglas. On the fifth day of hospitalization, the patient was discharged with a substantial improvement in abdominal pain. Five days after the discharge, her eosinophil count was elevated, and parasitic disease was therefore suspected. Anti-*Anisakis* IgG/A and IgE (RAST) antibody levels were elevated, confirming the diagnosis of intestinal anisakiasis. A review of 51 reported cases of intestinal anisakiasis suggests that the presence of ascites and measurement of anti-*Anisakis* antibody titers are helpful for diagnosis in cases presenting with nonspecific abdominal symptoms after consumption of raw or undercooked fish.

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Key words: abdominal pain, ascites, intestinal anisakiasis, raw fish

Introduction

Anisakiasis is a parasitic infection of the gastrointestinal tract caused by *Anisakis* larvae invading the gastrointestinal mucosa after ingestion of raw fish contaminated with the parasitic *Anisakis* spp¹. Approximately 20,000 cases of anisakiasis are reported worldwide annually, and more than 90% of these patients are from Japan². With the recent increase in the popularity of Japanese cuisine, such as sushi and sashimi, the incidence of anisakiasis has increased even outside Japan^{3,4}.

Although most anisakiasis cases are limited to gastric involvement, intestinal anisakiasis does rarely occur⁵. Intestinal anisakiasis can present with nonspecific abdominal manifestations, including abdominal pain, nausea, vomiting, and diarrhea⁶. Common complications include intestinal obstruction, perforation, and bleeding⁷. Typical findings of intestinal anisakiasis on computed tomogra-

phy (CT) include intestinal wall thickening with luminal narrowing, ascites, and fluid collection in the distal segment of the constricted small intestine^{8–10}. Its rarity and the nonspecificity of symptoms and CT findings make the diagnosis of intestinal anisakiasis difficult.

Here, we report a case of ascites of unknown etiology that was ultimately diagnosed as intestinal anisakiasis. We also reviewed the relevant literature on intestinal anisakiasis.

Case Report

A 63-year-old Japanese woman presented to the emergency department with progressive acute-onset abdominal pain and vomiting. She had consumed Mackerel pike (Sanma) sashimi 5 days prior to the visit. She denied having diarrhea, constipation, fever, shortness of breath, recent travel, or contact with sick individuals. The rest of

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review of systems was negative. Her vital signs showed a temperature of 36.2°C, pulse rate of 83 beats per minute, respiratory rate of 28 respirations per minute, blood pressure of 129/78 mmHg, and an oxygenation saturation of 99% on room air. Upon physical examination, she was alert and responsive, with a Glasgow Coma Scale score of 15 and a Japanese Coma Scale score of 0. Her respiration was unlabored, and her lungs were clear on bilateral auscultation. Cardiovascular examination results were unremarkable, with normal heart sounds and no murmurs, rubs, or gallops. Her abdomen was soft and non-distended; however, she had marked tenderness on palpation in the right lower quadrant, without guarding or rebound tenderness. Bowel sounds were normal. The remaining physical examination results were normal. Laboratory results revealed a white blood cell count of $9,200/\text{mm}^3$ (granulocytes, 86.8%; monocytes, 2.8%; eosinophils, 0.6%), hemoglobin level of 17.3 g/dL, and platelet count of $175,000/\text{mm}^3$. Her comprehensive metabolic panel and amylase level were within normal limits. Her HbA1c was 5.7%. An abdominal radiography revealed significant bowel gas. A chest radiography was unremarkable. Abdominal CT revealed thickened small bowel loops and ascites on the liver surface (Fig. 1). However, ischemic colitis could not be excluded. Thus, she was admitted for further evaluation and management of abdominal pain and ascites of unknown cause.

The initial differential diagnoses included a wide range of conditions from acute surgical abdomen to infectious enteritis. Infusion of lactate Ringer's solution and intravenous administration of cefmetazole 1 g twice a daily were initiated. Her symptoms improved rapidly; however, the presence of ascites could not be attributed to ordinary infectious enteritis. On the second day of hospitalization, contrast-enhanced abdominal CT scan demonstrated ileal wall thickening without poor contrast enhancement (Fig. 2). The ascites had moved from the liver surface to the pouch of Douglas while ischemic colitis was excluded. Her abdominal pain and serial abdominal examination had improved. On the fifth day of hospitalization, the patient was discharged with significant improvement of symptoms that were presumably due to common viral enteritis.

Five days after discharge, her complete blood cell count revealed an elevated the eosinophil count of 12.8%, which raised the suspicion of parasitic diseases. Anti-*Anisakis* IgG/A and IgE (RAST) antibody levels were elevated to 2.20 (normal value, ≤ 1.50) and 39.5 UA/mL (normal value, ≤ 0.34), respectively, which confirmed the

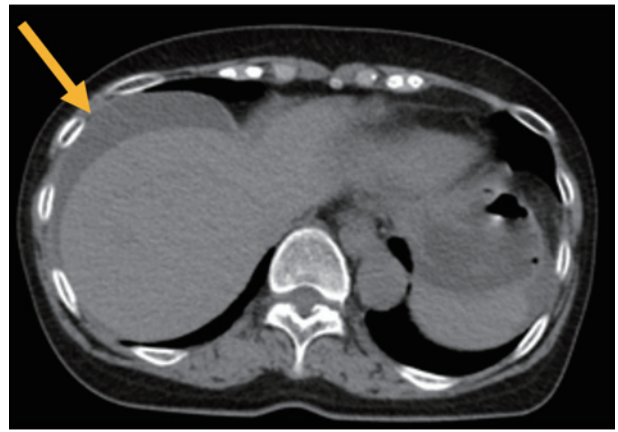


Fig. 1 Abdominal computed tomography showing ascites (arrow) on the liver surface.

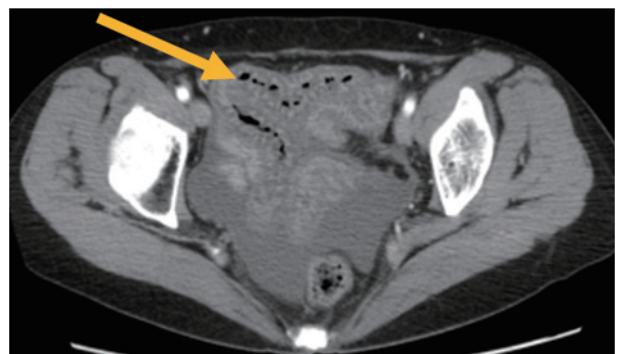


Fig. 2 Abdominal computed tomography with contrast enhancement demonstrating ileal wall thickening without poor contrast enhancement (arrow), as well as ascites extending from the liver surface to the pouch of Douglas.

diagnosis of intestinal anisakiasis.

Informed consent was verbally obtained from the patient.

Discussion

Here, we describe the case of a patient with ascites of unknown cause who was diagnosed with intestinal anisakiasis based on a history of raw fish consumption and elevated titers of anti-*Anisakis* antibodies. According to Ishikura et al.¹¹, gastric anisakiasis accounts for 95.6% of all cases, intestinal anisakiasis for 4.1% of cases, and infestation at other sites for 0.3%. Symptoms typically develop 12-24 hours after the ingestion of infected foods in cases of gastric anisakiasis, and 5-7 days after ingestion in cases of intestinal anisakiasis^{3,12,13}.

Intestinal anisakiasis can present with a broad range of abdominal symptoms, such as abdominal pain, nausea, vomiting, and diarrhea⁶. One study reported that complications include intestinal obstruction (50.7%), perforation

Table 1 Summary of our case and review of previously reported intestinal anisakiasis cases

Case	Age/ Sex	Presentation	Ascites (CT)	Endoscopy	Antibody	Operation	Author
1	63/F	AP, V	Yes	N/A	IgG/A Positive	N/A	Furuta et al. (Our case)
2	59/M	AP	No	Yes	N/A	Colectomy	Choi et al., 2019 ²⁶
3	30/M	AP, V	No	N/A	N/A	SB resection	Carlin et al., 2018 ²⁷
4	59/F	AP, V	Yes	Yes	N/A	N/A	Fujikawa et al., 2018 ²⁸
5	42/M	AP	Yes	N/A	N/A	SB resection	Shimamura et al., 2016 ¹⁸
6	47/M	AP, V	No	Yes	Antigen positive	N/A	Baeza-Trinidad et al., 2015 ²⁹
7	20/M	AP, V	Unknown	Yes	N/A	SB resection	Baron et al., 2014 ³⁰
8	52/F	AP, V, D	Yes	Yes	N/A	SB resection	Baron et al., 2014 ³⁰
9	36/F	AP	Unknown	Yes	N/A	SB resection & colectomy	Baron et al., 2014 ³⁰
10	39/M	AP	Unknown	Yes	N/A	SB resection	Baron et al., 2014 ³⁰
11	27/M	AP	Yes	Yes	N/A	SB resection	Baron et al., 2014 ³⁰
12	50/M	AP	Yes	Yes	N/A	SB resection	Baron et al., 2014 ³⁰
13	64/M	AP, N, D	Yes	N/A	N/A	SB resection	Shweiki et al., 2014 ¹²
14	62/M	AP	No	N/A	IgG/A Positive	N/A	Shrestha et al., 2014 ³¹
15	38/M	AP	Yes	N/A	IgG/A Positive	N/A	Shrestha et al., 2014 ³¹
16	47/F	AP	Unknown	N/A	IgG/A Positive	N/A	Shrestha et al., 2014 ³¹
17	63/F	AP, V	Yes	N/A	IgG/A Positive	N/A	Takano et al., 2013 ¹⁵
18	61/M	AP	Yes	N/A	N/A	SB resection	Kojima et al., 2013 ³²
19	14/M	AP, V	Unknown	N/A	N/A	SB resection	Juric et al., 2013 ³³
20	34/M	AP	Yes	N/A	IgG/A&E Positive	SB resection	Fujioka et al., 2012 ³⁴
21	30/M	AP	Yes	N/A	N/A	SB resection	Oshima et al., 2011 ³⁵
22	60/F	AP, V, FV	Unknown	N/A	N/A	SB resection	Kang et al., 2010 ³⁶
23	47/M	AP	Yes	Yes	Positive	SB resection	Fujii et al., 2009 ³⁷
24	51/F	AP, V	Yes	N/A	N/A	SB resection	Masui et al., 2006 ³⁸
25	59/M	AP, N	Yes	Only Gastrografin	IgG Positive	SB resection	Matsuo et al., 2006 ³⁹
26	62/M	AP, V	Yes	Only Gastrografin	IgE Positive	SB resection	Matsuo et al., 2006 ³⁹
27	32/F	AP, N, C	Yes	N/A	N/A	SB resection	Ramos et al., 2005 ⁴⁰
28	34/M	U, SOB, facial angioedema	Yes	N/A	N/A	SB resection	Ramos et al., 2005 ⁴⁰
29	80/M	AP, C	Unknown	N/A	N/A	SB resection	Ramos et al., 2005 ⁴⁰
30	49/M	AP, V	Yes	Yes	N/A	N/A	Ramos et al., 2005 ⁴⁰
31	76/M	AP, N	Yes	N/A	N/A	N/A	Ramos et al., 2005 ⁴⁰
32	53/F	AP	Unknown	N/A	N/A	N/A	Ramos et al., 2005 ⁴⁰
33	62/M	AP, N, C	Unknown	N/A	N/A	N/A	Ramos et al., 2005 ⁴⁰
34	25/M	AP, N, U	Unknown	N/A	N/A	N/A	Ramos et al., 2005 ⁴⁰
35	54/M	AP	Yes	N/A	N/A	SB resection	Yoon et al., 2004 ⁹
36	31/F	AP, V, D	Yes	N/A	N/A	SB resection	Caramello et al., 2003 ²⁴
37	50/M	AP, N, C	Yes	N/A	N/A	SB resection	Couture et al., 2003 ⁴¹
38	59/M	AP, V	Yes	N/A	N/A	SB resection	Sasaki et al., 2003 ¹⁴
39	43/M	AP	Yes	Yes	N/A	SB resection	Maggi et al., 2000 ⁴²
40	60/F	AP, V	Yes	N/A	N/A	SB resection	Takabe et al., 1998 ⁴
41	53/F	AP	Yes	N/A	Negative	SB resection	Kano et al., 1990 ⁴³
42	38/M	AP	Yes	N/A	Positive	SB resection	Kano et al., 1990 ⁴³
43	43/M	AP	Yes	N/A	Positive	N/A	Kano et al., 1990 ⁴³
44	52/M	AP	Yes	N/A	Positive	N/A	Kano et al., 1990 ⁴³
45	56/M	AP	Yes	N/A	Negative	N/A	Kano et al., 1990 ⁴³
46	33/M	AP	Yes	N/A	Positive	N/A	Kano et al., 1990 ⁴³
47	11/F	AP	Yes	N/A	Negative	Only exploratory laparotomy	Kano et al., 1990 ⁴³
48	30/M	AP	Yes	N/A	Positive	N/A	Kano et al., 1990 ⁴³
49	15/M	AP, V	Yes	N/A	Positive	N/A	Kano et al., 1990 ⁴³
50	35/F	AP, V	Yes	N/A	Positive	N/A	Kano et al., 1990 ⁴³
51	30/F	AP	Yes	Yes	N/A	SB resection	Appleby et al., 1982 ⁴⁴

AP, abdominal pain; C, constipation; CT, computed tomography; D, diarrhea; F, female; FV, fever; M, male; N, nausea; N/A, not applicable; SB, small bowel; SOB, shortness of breath; U, urticaria; V, vomiting

(8%), and bleeding (2%)⁷. The definitive diagnosis of intestinal anisakiasis is clinically challenging not only because there are no established diagnostic criteria, but also because symptoms and laboratory/imaging findings are nonspecific¹⁴. Therefore, obtaining a history of recent raw fish consumption is vital for accurate diagnosis^{6,15}. The differential diagnoses are broad and include acute appendicitis, ileitis, diverticulitis, cholecystitis, inflammatory bowel disease, peptic ulcers, small bowel obstruction, and intussusception^{16,17}. CT is useful in the diagnosis of intestinal anisakiasis^{8,9}, revealing marked submucosal edema and thickening of the intestinal wall, luminal narrowing, ascites, and/or fluid collection in the distal segment of the constricted small intestine⁸⁻¹⁰.

A review of 50 previously reported cases of intestinal anisakiasis, in addition to our case, is summarized in **Table 1**. The median age at diagnosis of intestinal anisakiasis was 47 years (range, 11-80 years). Of the 51 patients, 35 (69%) were men, and 16 (31%) were women. The most common presenting symptom was abdominal pain (n = 50, 98%), followed by vomiting (n = 17, 33%), diarrhea (n = 4, 8%), constipation (n = 4, 8%), urticaria (n = 2, 4%), and fever (n = 1, 2%). Our patient presented with abdominal pain, nausea, and vomiting, which are the most typical symptoms of intestinal anisakiasis.

While ascites was demonstrated on abdominal CT in 37 patients (72%), ascites was not found in four patients (8%), while ascites was not mentioned in 10 case reports (20%). This suggests that the presence of ascites has not typically been regarded as important as intestinal wall thickening for the CT diagnosis of intestinal anisakiasis. Shimamura et al.¹⁸ suggested four criteria for the diagnosis of intestinal anisakiasis: clinical features consistent with intestinal anisakiasis, a history of consumption of raw or undercooked fish within the previous 2 weeks, elevated levels of anti-*Anisakis* IgG/A or IgE antibodies, and presence of segmental intestinal edema and a distended proximal small bowel on CT. In fact, the clinical presentations of our patient met all four of these suggested criteria. However, the criteria do not include the presence of ascites. Nevertheless, our study of the literature indicates that more than 70% of cases present with ascites, and we propose that ascites should be considered a valuable diagnostic imaging index. In this context, our patient's clinical scenario, which included the above four characteristics plus ascites, appears to be very typical in intestinal anisakiasis.

Endoscopic evaluations were performed in 13 cases (25%), and an upper gastrointestinal series using Gastro-

grafin contrast agent was conducted in two cases (4%). Surgical intervention was performed in 33 patients (61%). In 37 cases with ascites on CT, 24 (65%) underwent surgical interventions, including exploratory laparotomy, small bowel resection, or colectomy, which suggests that more than 60% of cases of intestinal anisakiasis could mimic an acute surgical abdomen. Exploratory laparotomy was performed in 30 patients (60%). In these cases, intestinal anisakiasis was not initially suspected, but was revealed by exploratory laparotomy.

Anti-*Anisakis* antibodies are useful for the diagnosis of anisakiasis. Anti-*Anisakis* antibodies were measured in 19 patients (37%), of which 16 (31%) were positive. In contrast, antibodies were not measured in 31 patients (61%). An antigen was tested instead of an antibody in one case (2%). Elevated IgE antibody titers in the blood have a sensitivity of 100%, but a specificity of only 50% because of cross-reactions from other parasites, such as *Ascaris*, *Toxocara*, and *Echinococcus*^{19,20}, while elevated IgG/A antibody titers have a sensitivity of 70% and a specificity of 87%^{21,22}. In our case, anti-*Anisakis* antibodies were the diagnostic key; however, an earlier consideration of intestinal anisakiasis as a differential diagnosis of ascites might have prompted earlier testing for anti-*Anisakis* antibodies. Although paired sera were not obtained in our case, the use of such paired sera could provide more accurate diagnostic yield by mitigating the potential for false negatives during the early stages of symptom onset²¹.

Based on the above findings, we believe that the presence of ascites and the measurement of anti-*Anisakis* antibody titers may be useful for the diagnosis of intestinal anisakiasis. It is reasonable to propose that intestinal anisakiasis should be included on the list of differential diagnoses, especially when patients present with nonspecific abdominal symptoms after eating raw or undercooked fish^{6,23,24}. According to Ido et al.²⁵, the actual number of intestinal anisakiasis cases is estimated to be greater than previously reported. Understanding the importance of ascites and anti-*Anisakis* antibodies will hopefully lead to the detection of more undiagnosed cases of intestinal anisakiasis.

Conclusions

Herein, we report a case of intestinal anisakiasis associated with ascites. It is suggested that intestinal anisakiasis should be considered in patients with nonspecific abdominal symptoms accompanied by the presence of ascites on abdominal CT scan, particularly in patients with a recent history of recent raw fish consumption, and that

the measurement of anti-*Anisakis* antibodies is beneficial for diagnosis.

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