

A Tubo-Ovarian Abscess Caused by *Salmonella enterica*

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Abstract

A tubo-ovarian abscess (TOA) is a serious complication of pelvic inflammatory disease (PID), which is usually associated with polymicrobial infections, with a predominance of anaerobic bacteria. *Salmonella* species are rarely associated with TOA, as only 11 cases have been documented in the literature. We present the case of an 18-year-old virgin who contracted a *Salmonella enterica* (*S. enterica*)-induced TOA. After experiencing an episode of gastroenteritis 13 days earlier, attributed to chicken consumption, the patient revealed severe abdominal pain, fever, diarrhea and vomiting. A right adnexal mass was found by imaging, and symptoms continued even after starting broad-spectrum antibiotic treatment. For this reason, an exploratory laparotomy was considered crucial, and it confirmed a right TOA, necessitating surgical management, including a right salpingectomy and abscess drainage. Postoperative cultures identified *S. enterica* as the causative organism. The patient recovered uneventfully after targeted antibiotic therapy. A *Salmonella*-associated TOA was observed in a non-sexually active patient, with the most probable route of transmission being an ascending infection following recent gastroenteritis. No underlying gynecological pathology was found. Since unspecific symptoms of TOA might imitate those of other acute abdominal disorders, this article emphasizes the difficulties in diagnosing this pathology in sexually inactive patients. Including rare pathogens such as *Salmonella* species in the differential diagnosis of TOA is essential. Early recognition, combined with appropriate surgical treatment, is critical for a successful outcome if conservative management fails.

Keywords: Tubo-ovarian abscess; *Salmonella*; Case report; Virgin; Exploratory laparotomy

Introduction

Tubo-ovarian abscess (TOA) is a complex infectious adnexal mass that typically arises as a complication of pelvic inflammatory disease (PID). These abscesses are often polymicrobial, with anaerobic bacteria being the predominant pathogens [1].

While *Salmonella* is a gram-negative bacterium widely known as a major cause of gastroenteritis worldwide, its involvement in TOA is exceedingly rare [2, 3]. To date, only 11 cases of *Salmonella*-associated TOA have been documented in the literature [3-13]. This article aims to present the case of an 18-year-old virgin diagnosed with a TOA caused by *Salmonella enterica* (*S. enterica*). The case is analyzed in detail, hence providing insights into the clinical presentation, diagnosis and management of this rare condition.

Case Report

Investigations

An 18-year-old female presented to our Emergency Department with acute onset of abdominal pain, accompanied by four episodes of vomiting, one episode of diarrhea, and a single episode of fever with chills within 24 h prior to admission. Her medical history included a recent episode of gastroenteritis 13 days prior to admission, attributed to chicken consumption. The symptoms, consisting of two episodes of diarrhea, one episode of vomiting, and dizziness, lasted for 1 day and resolved quickly after self-treatment with loperamide, famotidine and domperidone. On our clinical examination, the patient exhibited a mildly rigid abdomen with diffuse tenderness, a positive Murphy's sign, and rebound tenderness. A vaginal examination was deferred as the patient reported virginity.

Diagnosis

Her vital signs were stable, and she had an elevated body tem-

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Figure 1. Axial post-contrast computed tomography (CT) scan of the abdomen shows a cystic lesion in the right adnexa (arrow) with a thick enhancing wall. Fat stranding and free fluid are also present (arrow-heads).

perature of 38.6 °C. Initial laboratory investigations revealed leukocytosis with a white blood cell (WBC) count of 21,560/ μ L and an elevated C-reactive protein (CRP) level of 134 mg/L. An urgent abdominal ultrasound revealed a distended appendix measuring 7.4 mm in diameter. Gynecological ultrasonography identified a cystic adnexal mass measuring 67 \times 61 \times 47 mm, characterized by mixed echogenicity. To further evaluate the findings, an emergency contrast-enhanced computed tomography (CT) scan of the abdomen and pelvis was performed. CT of the lower abdomen revealed a cystic hypodense lesion on the right side of the pelvis with a thick enhancing wall. Fat stranding and free fluid were also present (Fig. 1). These imaging features were consistent with PID, a right adnexal lesion, and secondary inflammatory changes involving the appendix, small bowel, and colon. Additionally, vaginal fluid and blood cultures were obtained.

Treatment

The patient was admitted to the hospital and initiated on broad-spectrum antibiotic therapy consisting of cefoxitin 2 g three times daily intravenously (IV), metronidazole 500 mg twice daily IV, and doxycycline 100 mg twice daily orally. Painkillers, antiemetics and antipyretics were administered as needed, using tramadol, ondansetron and paracetamol. During hospitalization, the patient remained hemodynamically stable. Her clinical course was characterized by persistent abdominal pain localized to the epigastric region, periumbilical area, and right hypochondrium, initially intense and later becoming milder. She also experienced two episodes of diarrhea that lasted 2 days, one to three episodes of vomiting per

day, and a fever once or twice daily, ranging from 38 °C to 39 °C, which was alleviated with paracetamol. A daily complete blood count and biochemical tests, including electrolytes, coagulation profiles and inflammatory markers, were performed almost daily throughout the hospitalization. The results revealed elevated inflammatory markers, with an initial CRP of 277 mg/dL and an increase in WBCs, which later showed slight improvement with the administration of antibiotics. For diagnostic workup and laboratory evaluation, a panel of tests was conducted, including severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) testing, serum human chorionic gonadotropin (β -HCG), thyroid function tests, upright abdominal radiography, upper and lower abdominal ultrasonography, urinalysis, B-Koch urine culture, stool parasitological examination, stool culture for *Clostridium* species pluralis (spp.), and stool analysis for the detection of toxins A, B and C. All cultures were sterile, and no other pathological findings were observed. Due to mild inflammatory marker elevation and persistent symptoms despite antibiotics, cefoxitin was discontinued, and ceftriaxone 2 g IV daily was initiated, along with the previous antibiotic regimens. Blood cultures obtained at the Emergency Department were negative, and vaginal fluid cultures revealed the growth of *Gardnerella vaginalis*. Following the modification in antibiotic therapy, the patient showed no significant clinical improvement, with persistent one to two episodes of vomiting per day and febrile episodes once or twice daily, ranging between 38 °C and 38.6 °C. Diagnostic evaluation continued with a repeat gynecological ultrasound, which revealed no new findings or improvement in the pre-existing condition. Urine cultures were performed using Ziehl-Neelsen staining, auramine-rhodamine, BAC-TEC MGIT, Loewenstein-Jensen, and Myco/F Lytic bottle methods. An interferon-gamma release assay blood test was also conducted, alongside serological testing for hepatitis B virus (HBV), hepatitis C virus (HCV), human immunodeficiency virus (HIV) and venereal disease research laboratory (VDRL). All tests returned negative for detectable pathogens. Given the ongoing febrile episodes, the antibiotic regimen was altered to Tazocin 4.5 g four times a day, and doxycycline was continued. Subsequently, a magnetic resonance imaging (MRI) of the abdomen and pelvis was performed. The imaging clearly depicted the complex cystic lesion, with heterogeneous signal intensity on T1 and T2 sequences, displacing the right adnexa inferiorly. The lesion showed areas of restricted diffusion, high signal in diffusion-weighted imaging (DWI), and low signal in an apparent diffusion coefficient (ADC) map, findings concomitant with abscess formation (Fig. 2). Due to persistent symptoms, blood cultures were obtained, which were negative. Following a surgical consultation, the patient underwent an exploratory laparotomy. Prior to surgery, she was under antibiotic treatment with Tazocin 4.5 g four times daily IV and doxycycline 100 mg twice daily orally. She also received 2,500 IU of bempiparin and 40 mg of omeprazole IV before the procedure. The patient was kept *nil per os* for 1 day prior to surgery. Intraoperatively, a right TOA was identified, for which a right salpingectomy and removal of the abscess wall were performed. Associated pelvic fluid collections were drained, and specimens were sent for comprehensive microbiological and molecular analysis, including cultures for bacte-

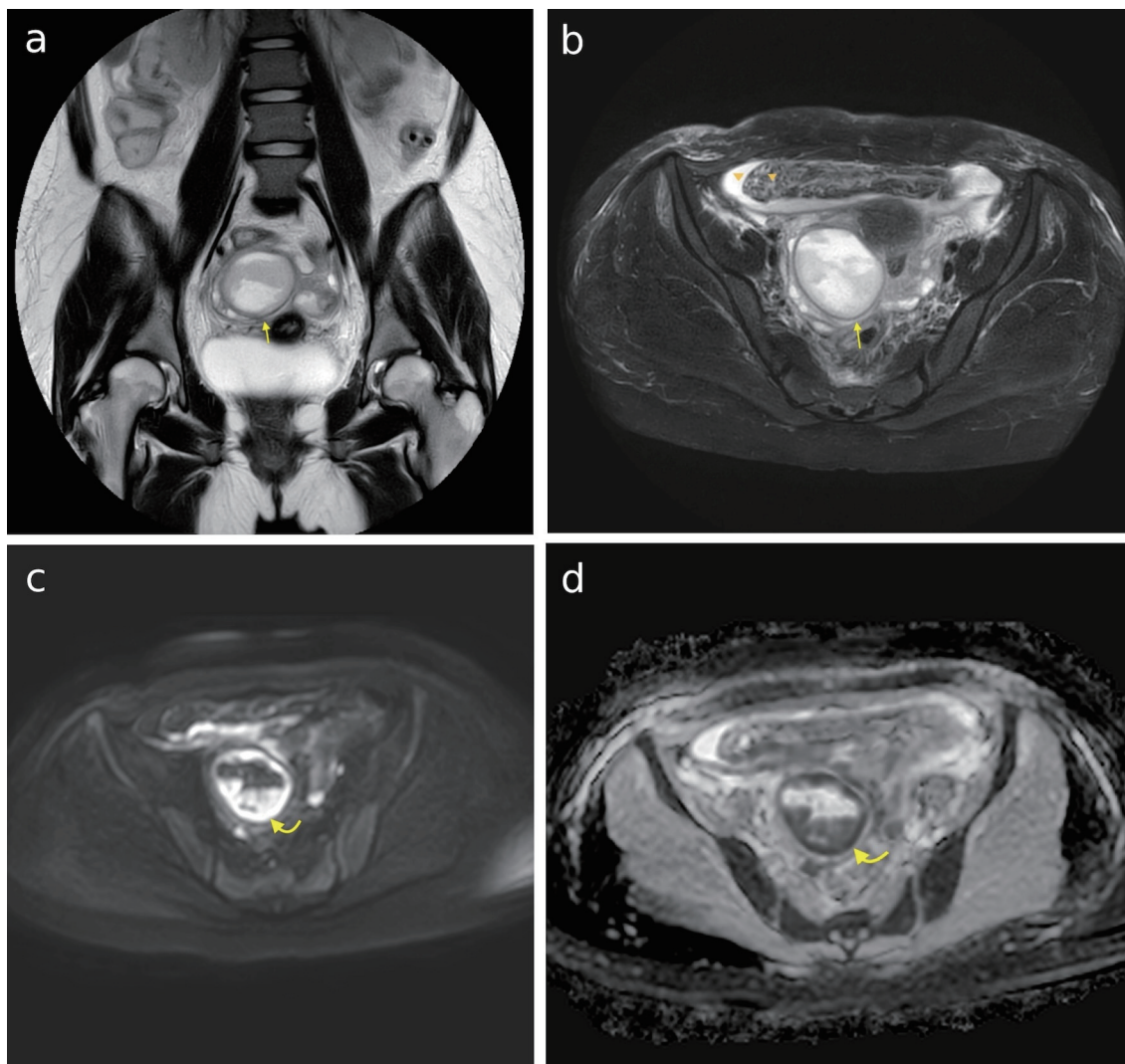


Figure 2. (a) Coronal T2-weighted image of the abdomen shows a highly heterogeneous complex cystic lesion in the right side of the pelvis (arrow). (b) Axial T2-weighted fat-saturated image shows a large complex cystic lesion in the right side of the pelvis (arrow), displacing the right adnexa inferiorly. Fat stranding and free fluid are also present (arrowheads). (c) Axial DWI image at $b = 1,000 \text{ s/mm}^2$, showing areas of high signal intensity within the lesion (curved arrow). (d) Same areas depict low signal in the ADC map corresponding to areas of restricted diffusion (curved arrow). DWI: diffusion-weighted imaging; ADC: apparent diffusion coefficient.

rial pathogens, mycobacteria, fungi, and sexually transmitted infections (STIs).

Follow-up and outcomes

Postoperatively, the patient remained hemodynamically stable and afebrile, with an uneventful recovery. Cytological examination of the pelvic abscess fluid and peritoneal fluid revealed findings consistent with acute inflammation, with no evidence of malignancy. Histological analysis of tissue samples from the pelvic abscess, right fallopian tube, and the wall of the ovarian abscess demonstrated acute and chronic inflammation, with no evidence of malignancy. Molecular investigation was performed on cytological material obtained from

the pelvic abscess fluid using real-time multiplex polymerase chain reaction (PCR), targeting STI markers such as *Ureaplasma parvum*, *Ureaplasma urealyticum*, *Mycoplasma hominis*, *Mycoplasma genitalium*, *Trichomonas vaginalis*, *Neisseria gonorrhoeae*, *Chlamydia trachomatis*, and *Candida* species, which returned negative for tested pathogens. Peritoneal fluid cultures were also conducted using Gram staining, aerobic and anaerobic cultures, as well as agar cultures for fungi, all of which returned negative results. Pus cultures were performed for anaerobic, fungal, and mycobacterial organisms, all of which yielded negative results. However, aerobic culture on MacConkey agar identified *Salmonella* spp. Further, PCR and serotyping using slide agglutination were obtained to reveal *S. enterica* subsp. *enterica* serovar *Enteritidis*. Antimicrobial susceptibility testing was conducted on the *S. enteritidis* iso-

late from the patient's clinical sample using the disk diffusion technique to determine the susceptibility to a panel of antibiotics. The isolate was susceptible to all antibiotics tested, with minimum inhibitory concentrations (MICs) well below the resistance threshold. Following infectious disease consultation, the patient was prescribed ceftriaxone 2 g IV for 7 days, followed by oral ciprofloxacin 600 mg twice daily and metronidazole 500 mg three times daily for 7 days. A follow-up evaluation was scheduled for 10 days post-treatment, and it was uneventful.

Discussion

The presented case adds to the literature on TOA due to *Salmonella*, with 11 similar cases identified through a review using the keywords "tubo-ovarian abscess" and "*Salmonella*" [3-13]. In this report, we describe a case of a right TOA caused by *S. enterica* in an 18-year-old female with no history of sexual intercourse. To the best of our knowledge, this is the second such case involving a sexually inactive patient, the first being a 42-year-old female reported by Yakupogullari et al in 2019 [13].

Potential routes of transmission include lymphatic or hematogenous spread following bacteremia, ascending infection from the lower genital tract, or local spread from an infected adjacent organ such as the appendix or small intestine [3, 7, 13]. Regarding our patient, the most likely route of transmission must be through ascending infection from contaminated stool, following the episodes of diarrhea 13 days prior to admission, attributed to chicken consumption.

According to previously reported cases, most of the patients had an underlying gynecologic condition, suggesting that pre-existing pathology predisposes to localized *Salmonella* infection, either known prior to diagnosis or incidentally discovered during evaluation [14]. Endometrioma and endometriotic cysts were the most common underlying conditions, reported in five cases of TOA due to *Salmonella* by Ghose et al [4], Kostiala et al [5], Thaneemalai et al [6], Manning et al [8], Kudesia et al [10]. Additionally, Selvam et al [12] reported one case associated with a pre-existing dermoid cyst [12]. It should be noted that patients with systemic lupus erythematosus (SLE) or immunocompromised patients can present with an increased risk of developing localized infections following invasive salmonellosis [11, 14, 15]. Furthermore, there has been a case of adnexal abscess in a female patient who was a potential gastrointestinal *Salmonella* carrier following *in vitro* fertilization (IVF). In that case, it was hypothesized that the oocyte collection needle was contaminated during a possible stick injury to the bowel [13]. Notably, in our patient and the 47-year-old female reported by Valayatham et al [9], no visible ovarian abnormalities or pre-existing disease were identified.

TOA in young, sexually inactive girls poses a diagnostic challenge. The rarity of *Salmonella*-induced TOAs often leads to a lower index of suspicion, contributing to diagnostic delays [3, 16]. As noted in previous cases, this rare condition can mimic ovarian cysts, tumors, or acute abdomen [12, 14-17]. In our case, the patient's presentation with nonspecific symptoms such as abdominal pain, fever and gastrointestinal dis-

turbances, coupled with the absence of typical risk factors for PID, further obscured the clinical picture. In the current case, it was only upon surgical exploration and subsequent microbiological analysis that *Salmonella* was identified as the causative pathogen. This underscores the importance for clinicians to maintain a broad differential diagnosis when evaluating pelvic masses, especially in patients presenting with gastrointestinal symptoms, and to consider uncommon pathogens like *Salmonella* within the etiological spectrum of TOAs [15]. Laboratory diagnosis of *Salmonella*-induced TOAs primarily relies on the isolation of the organism from abscess fluid or tissue samples. Culture techniques remain the gold standard, with growth typically observed on selective media such as MacConkey agar. In our case, pelvic fluid collections were aspirated, and samples were submitted for comprehensive microbiological and molecular testing, including cultures for bacterial pathogens, mycobacteria, fungi, and STIs. Histopathological examination might reveal features such as hemorrhagic and degenerative changes within the cyst wall, often associated with endometriotic tissue, as noted in similar case reports. Serological tests and stool cultures can provide valuable information regarding concurrent or preceding enteric infections, providing additional context for the diagnosis [3].

In our case, broad-spectrum antibiotic therapy was initiated, comprising cefoxitin, metronidazole, and doxycycline. Successful conservative management with antibiotics alone has not been described in the literature. Surgical removal of the infected abscess has been deemed necessary for all previously reported cases of TOA due to *Salmonella* [3-13]. All 11 reported cases of TOA due to *Salmonella* have been treated after laparotomy or laparoscopy, including our case. However, while antibiotics remain the first-line treatment for TOAs, unsuccessful cases should proceed to drainage under ultrasound guidance or surgical intervention [12]. Empirical broad-spectrum antibiotic therapy should be initiated promptly, targeting both typical pelvic pathogens and *Salmonella* species. Following culture and sensitivity results, antibiotic regimens can be tailored accordingly. The most widely used and effective protocols include a combination of clindamycin, gentamicin, and ampicillin; cephalosporin with gentamicin and metronidazole; or cephalosporin combined with metronidazole. Surgical management is considered in patients who do not respond to antibiotic therapy within 72 h, present with large abscesses, or exhibit signs of rupture or sepsis. Procedures range from minimally invasive techniques, such as image-guided percutaneous drainage, to more extensive surgeries like laparoscopic or open salpingo-oophorectomy. The type of procedure depends on factors such as abscess size, patient stability, and reproductive considerations. Early and aggressive treatment is crucial to prevent complications such as abscess rupture, peritonitis, and sepsis, thereby preserving reproductive function and reducing morbidity [17].

Learning points

The diagnostic complexity of TOA caused by *Salmonella* lies in its ability to mimic other acute abdominal pathologies, such as ovarian cysts, tumors, or appendicitis, particularly in pa-

tients without a history of sexual activity or pre-existing gynecologic conditions. Our findings emphasize the importance of considering *Salmonella* as a potential causative pathogen in TOAs, even in atypical clinical presentations. Despite the use of broad-spectrum antibiotics as initial therapy, surgical intervention remains a cornerstone of effective treatment, as demonstrated by previously reported cases, including this one. This case highlights the need for heightened clinical suspicion, thorough history-taking, and a multidisciplinary approach for timely diagnosis and management of this rare but significant condition.

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None to declare.

Financial Disclosure

None to declare.

Conflict of Interest

None to declare.

Informed Consent

Informed consent was obtained.

Author Contributions

NK, MK and SM observed the patient. EE performed a radiology consultation. PP observed the patient and performed medical treatment. NK, MK and SM drafted, reviewed and edited the manuscript. PP and NM reviewed and edited the manuscript. All authors have approved the final article for journal publication.

Data Availability

Any inquiries regarding supporting data availability of this study should be directed to the corresponding author.

Abbreviations

TOA: tubo-ovarian abscess; PID: pelvic inflammatory disease; STIs: sexually transmitted infections; WBC: white blood cell; CRP: C-reactive protein; IV: intravenously; IVF: *in vitro* fertilization; CT: computed tomography; MRI: magnetic resonance imaging; DWI: diffusion-weighted imaging; ADC: apparent diffusion coefficient; MIC: minimum inhibitory concentration

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