INTRODUCTION

Actinomycosis is a rare chronic or subacute infectious disease resulting in suppurative and granulomatous inflammation. The disease is caused by anaerobic or facultatively anaerobic, acid-resistant Gram-positive bacteria of *Actinomyces* species. In the past classified as fungi, they derive their name from the shape of their colonies, whose branched filamentous forms resemble traveling rays. In healthy subjects, *Actinomyces* form part of normal flora of the oral cavity or less commonly, a part of the gastrointestinal tract and the genital tract. *Actinomyces* are thus rated among opportunistic microorganisms. Infections are endogenous and occur more commonly in males (M:F = 3:1), usually in subjects aged 20–50. The bacteria enter the body through the damaged mucous membrane or the skin, most commonly during tooth extractions, abdominal surgery or when having an intrauterine device (IUD) inserted. In locations with favorable growth conditions (anaerobic environment), *Actinomyces* provoke development of abscesses, having a tendency to form fistulas; they spread contiguously. In humans, actinomycosis is most commonly caused by *Actinomyces israelii*, *Actinomyces propionica* and *Actinomyces naeslundii*, although at least 39 species of this bacterium have been described. A distinctive feature of this type of inflammation is the presence of "actinomycosis grains", sulfur colonies in the form of 0.1–5 mm grains in purulent contents secreted from the fistula. A microscopic test along with the presence of distinctive macroscopic features of purulent discharge leads to a diagnosis, which should be confirmed by microbiological evaluation. Actinomycosis treatment involves surgical drainage of abscesses and the prolonged use of penicillins administered in high doses.

CASE REPORT

A 50-year-old female patient was admitted to the Department of Allergy and Immunology Jagiellonian University School of Medicine to diagnose a cause of subfebrile temperature and left-sided subcostal pain of 1.5 years’ duration. She also complained of hip joint, knee joint and spinal pains. Within 4 months preceding hospitalization she lost 4 kg. She did not report any urination or defecation disorders. The patient’s father died of stomach cancer at the age of 49. At 24, the patient had her gallbladder removed due to cholelithiasis. Physical examination revealed a mass of indistinct borders and a diameter of ~5–10 cm that was found in the left subcostal area. The location was slightly tender to palpation. Furthermore, on the skin of the left shank, a ~3 mm wide single ecchymosis was noticed, and the limb skin demonstrated livedo reticularis. Small ecchymoses on the limbs and the face had subsided spontaneously.

ABSTRACT

Actinomycosis is a rare infectious inflammatory disease caused by the bacteria of the *Actinomyces* species. Infection occurs following damage to the skin or mucous membrane. This report presents a case of a 50-year-old female patient with subfebrile temperature, weight loss and pain, who used to have an intrauterine device. Masses suggestive of an advanced cancer were detected in her pelvis and abdominal cavity. A diagnosis of actinomycosis was made after histopathological examination of the tissue sample.
Area getting wider towards the front and sides of the lesion. Contrast-enhanced computed tomography (CT) of the abdomen and the pelvis showed heterogeneous infiltration, 6 cm in diameter in the anatomical location of the right ovary, intraperitoneally, with strong enhancement. Moreover, below this infiltrate, a small, belt-shaped fluid collection was observed (Figure 1), and in the abdominal wall, within the left subcostal area, the patient demonstrated a 10 × 4 cm heterogeneous, strongly contrast enhancing infiltration (Figure 2). The fine-needle aspiration biopsy of the abdominal wall lesion was performed. Cytology of the biopsy sample, performed in the Chair of Pathomorphology Jagiellonian University School of Medicine, showed pus with bacterial colonies typical of actinomycosis (Figure 3).

After initiating antibiotic therapy using amoxicillin-clavulanic acid, the patient was transferred to the department of gynecology. On surgery, a tumorous infiltration located in the abdominal wall in the right inguinal area and a similar infiltration in the bladder wall, which was connected with the right-sided inflamed, adnexa were removed. Moreover, the uterus with bilateral adnexa was resected. Palpation and inspection of the abdomen showed no other pathological lesions. The infiltration in the left-sided subcostal area, described earlier in the CT and ultrasound examination, subsided after antibiotic therapy and the pain disappeared.

Histological examination of the bladder tumor lesion showed the presence of a mixed-cell, reactive inflammatory infiltration was reported, whereas the material obtained from the right ovary and adnexa showed scattered unspecific infiltrations with granulation centers of an “around a foreign body” type were found. Within the fallopian tubes, purulent exsudates were observed (Figure 4). The described histopathological lesions may correspond to the picture of lesions found in actinomycosis.

**DISCUSSION** Depending on the lesion location, several forms of actinomycosis, i.e., cervicofacial, thoracic, abdominal and pelvic, can be distinguished. The patient demonstrated a rare, mixed abdominopelvic form. Available data points out the association between use of IUD and occurrence of pelvic actinomycosis. This is based on two observations. First, the common presence of *Actinomyces* in cervical smears obtained from users of IUD as compared to those who do not use IUD. Second, the invasive pelvic actinomycosis occurs nearly exclusively in women using IUD. The patient presented here reported a history of several years’ use of IUD. It is highly probable that infection was transmitted through the genital tract. It seems that the original lesion was a tumorous infiltration of the right ovary mimicking the cancer, whereas infiltrations of the abdominal wall and the bladder could have been secondary. A few years earlier the patient had not undergone
any surgery within the abdominal cavity and abdominal actinomycosis is most commonly associated with surgical procedures.

In order to prevent invasive actinomycosis, users of IUDs, in whom *Actinomyces* were detected in the cervical smear, should be suggested to remove the intrauterine device, and those, who plan to insert an IUD, should be firmly advised against it. Examination of the cervical canal contents for the presence of *Actinomyces* in those patients could undoubtedly reduce the incidence of the invasive form of pelvic actinomycosis.

Actinomycotic lesions, with the concomitantly affected pelvis and abdominal cavity have rarely been described. Such manifestations are most commonly encountered in case of cancer located originally in the ovary. The described case indicates that in differential diagnosis of such lesions, except for cancer, an infection, like actinomycosis, should also be considered. In the latter case, even with highly advanced local lesions, prognosis is quite good on condition of an accurate diagnosis and the implementation of appropriate therapy.

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