Pyomyositis associated with chemotherapy for endometrial cancer: a case report

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Abstract
Pyomyositis is a rare complication of chemotherapy for non-hematological malignancies. A 58-year-old woman with endometrial carcinoma, in whom pyomyositis developed during adjuvant chemotherapy, was presented in this report. After initiating empirical antibiotic therapy for febrile neutropenia, screening CT showed multiple abscesses in the lower limbs. Operative drainage of the abscess was effective.

Keywords: Pyomyositis, Endometrial carcinoma, Chemotherapy

Background
Pyomyositis is originally known as an infectious disease of the large skeletal muscles, which is common in tropical areas and often results in abscess formation and sepsis. In patients with pyomyositis, *staphylococcus aureus* is detected more often than *streptococcus* and other bacteria. Important factors of pyomyositis are reported to be local mechanical trauma, parasitic infections, and malnutrition. Furthermore, it is recently reported that pyomyositis is associated with the immunocompromised host due to infection with human immunodeficiency virus (HIV), hematological malignancies, and neutropenia secondary to oncologic chemotherapy [1].

To date, pyomyositis developing as a complication of chemotherapy in patients with gynecologic malignancy has been rarely reported [2]. In this report, we presented a patient with pyomyositis who had received systemic chemotherapy for endometrial carcinoma.

Case presentation
A 58-year-old previously healthy Japanese woman (body mass index 22.1) presented sudden massive postmenopausal bleeding, and diagnosed endometrial carcinoma after gynecological, pathological, and radiological examinations. She did not have family history of cancer. She underwent total abdominal hysterectomy, bilateral salpingo-oophorectomy, pelvic lymphadenectomy, and sampling of para-aortic lymph nodes. Histopathology revealed endometrioid adenocarcinoma grade 1. Tumor cells involved the serosa of uterine corpus. The pathological diagnosis of endometrial carcinoma stage IIIc, T3a N1 (right obturator lymph node) M0, was made. The patient was planned to be given whole pelvic external irradiation followed by systemic chemotherapy.

External radiotherapy was delivered with an 18-MV photon beam for 5 days each week. Whole pelvic irradiation was delivered to a total dose of 45 Gy in 25 fractions. The first course of chemotherapy consisted of docetaxel (70 mg/m2) and carboplatin (AUC = 5) administrated on Day 0. On Day 11, she presented at our outpatient clinic with worsened general fatigue, appetite loss, and restlessness. Her body temperature developed to 39.1°C. Physical examination showed muscular weakness and proximal dominant lower limb muscle ache. Routine blood test revealed decreased white blood cell count (1.6 × 10^3 /μL), decreased platelet count (5.3 × 10^4 /μL), increased C-reactive protein (46.14 mg/dL (normal, 0 to 0.3)), increased serum CPK (8152 IU/L), increased LDH (638 IU/L), increased serum creatinine (2.12 mg/dl) and increased blood urea nitrogen (44.0 mg/dl). Imaging of her head and chest-abdomen with computed tomography showed no remarkable changes. Empirical anti-bacterial chemotherapy for pyrogenic leukocytopenia with cefepime (4 g/day) was started. She was found to have *Streptococcus dysgalactiae equisimilis* bacteremia. She received
intravenous antimicrobial treatment with vancomycin (2 g/day) and clindamycin (1.8 g/day); which improved her conditions. However, her body temperature developed to 39.7°C on Day 14 again, and prolonged fever elevation maintained at 38 to 39°C. A repeated physical examination showed tenderness on left medial thigh. CT scan demonstrated the formation of multiple abscesses in the both femoral muscles on Day 21 (Figure 1).

The patient underwent open surgery. The incision was made in the left medial thigh and opened the fascia. Further incision was made in the swelling medial vastus muscle. About 50 mL of pus was drained. A Penrose drain was left in the abscess and the incision was closed. The drained pus showed the same bacteria with the bacteremia. The fever decreased and general conditions improved. In addition, 7 weeks oral administration of clindamycin (1.2 g/day) and levofloxacin (0.5 g/day) suppressed the relapse of inflammation.

Because of these severe complications, an additional course of chemotherapy was not given. After 40 months of this event, this patient is still alive with the disease.

Discussion

Bacterial pyomyositis is a purulent infection of the large skeletal muscle, and is most commonly involved in the gluteal areas and the thigh; which results in abscess formation. It is an endemic disease in the tropics, and seems to occur only occasionally in the temperate zones. However, pyomyositis is increasingly recognized in non-tropical areas as one of the complications in the immunocompromised host associated with HIV infection, diabetes mellitus, and chemotherapy-induced neutropenia. Both malignant and non-malignant hematological diseases have been reported in association with pyomyositis [1,3]. A review paper reports 45 patients with hematological neoplastic diseases associated with pyomyositis [1]. Pyomyositis is an uncommon complication of chemotherapy for non-hematological malignancies. To our knowledge based on MEDLINE, only nine cases of solid tumor-related pyomyositis have been reported (Table 1). Among them, two have retroperitoneal malignant teratomas [4], two lung cancer [5,6], two breast cancer [3,7], one colon cancer [3], one glottic cancer [3], and one endometrial cancer [2].

Primary pyomyositis is believed to be caused by transient bacteremia, because it develops without an obvious penetrating injury or any other clear portals of entry in the majority of cases [3]. Development of pyomyositis in patients with neoplastic diseases after chemotherapy is usually ascribed to neutropenia and/or immunodeficiency caused by cancer. However, subclinical myopathy secondary to malignancy and/or its treatment is another possible cause [3]. Several

Figure 1 Multiple abscesses in the lower extremities showing low density areas with enhancement (arrows).
chemotherapeutic drugs including anthracyclines, vinca alkaloids, and corticosteroids have been reported to induce muscle toxicity-related pyomyositis [7,8]. It is recently reported that a woman with endometrial cancer develops pyomyositis after the first cycle of carboplatin and paclitaxel [2]. Our patient also received docetaxel and dexamethasone. However, the muscle toxicity of docetaxel has never been reported. Whether pyomyositis observed in the above two cases was related to the drug was unclear.

Diagnosis of pyomyositis is facilitated by ultrasonography, MRI, or CT scanning of the affected area, and by aspiration of fluid for microbiological testing. According to previous literature, Staphylococcus aureus is the most common cause of pyomyositis. Several other bacteria including Gram-negative organisms, other Gram-positive organisms than Staphylococcus aureus (predominantly Streptococcus), anaerobes, mycobacteria, and fungi are implicated as a cause of pyomyositis [3].

Early diagnosis enables complete drainage of purulent materials and successful treatments, and leads to resolution in the vast majority of cases. The immune status of the host, clinical courses, and the number of abscess materials and successful treatments, and leads to resolution in the vast majority of cases. The immune status of the host, clinical courses, and the number of abscesses should be considered when determining the length of the treatment for pyomyositis.

**Conclusion**

We described an unusual case of pyomyositis associated with chemotherapy for gynecologic malignancy. This should be kept in mind in the differential diagnosis of febrile neutropenia.

**Consent**

Informed consent was obtained from the patient for publication of this case report and accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

**Competing interests**

The authors declare that they have no competing interests.

**Author's contributions**

YN contributed mainly in designing, literature review, and writing work. MY, MH, and MN have operated this case. SN, SA, MY, and TI gave advices and edited the discussion. All authors read and approved the final manuscript.

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**Table 1 Review of patients with pyomyositis associated with solid malignancies**

| Patient (age/sex) | The origin of cancer | Chemotherapy | Site of pyomyositis | Surgical drainage | Bacteria | Reference |
|-------------------|----------------------|--------------|---------------------|------------------|----------|-----------|
| 1 43/M            | Retropertoneal teratoma | VCR, BLM, CDDP, MTX, VP-16, CPA | Psoas muscle | Done | Bacteroides fragilis | [4] |
| 2 40/M            | Retropertoneal teratoma | VCR, BLM, CDDP, MTX, VP-16, CPA | Psoas muscle | Done | Bacteroides fragilis | [4] |
| 3 67/M            | Lung                  | CDDP, VDS, MMC | Lower limb | Done | Staphylococcus aureus | [5] |
| 4 47/F            | Breast                | 5-FU, DTX, CPA | Carf               | Done | Staphylococcus aureus | [7] |
| 5 62/M            | Lung                  | AMR, VNR     | Upper arm, lower limb | Done | Staphylococcus aureus | [6] |
| 6 NR              | Breast                | Unknown      | NR                 | NR               | NR       | [3] |
| 7 NR              | Colon                 | Unknown      | NR                 | NR               | NR       | [3] |
| 8 NR              | Glottis               | Unknown      | NR                 | NR               | NR       | [3] |
| 9 58/F            | Endometrium           | CBDCA, PTX   | Gluteus medius     | CT-guided drainage | Staphylococcus aureus | [2] |
| 10 58/F           | Endometrium           | CBDCA, DTX   | Lower limb         | Done | Streptococcus dysgalactiae equisimilis | This case |

AMR Amrubicin; BLM Bleomycin; CBDCA Carboplatin; CDDP Cisplatin; CPA Cyclophosphamide; DTX Docetaxel; DXR Doxorubicin; MMC MitomycinC; MTX Methotrexate; NR Not recorded; PTX Paclitaxel; VNR Vinorelbine; VP-16 Etoposide; VCR Vincristine; VDS Vindesine.
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