Case report

A 45-year-old married woman presented with 2 months history of a vulval mass as large as a soybean. There was no swelling, pain, vulval itching or causalgia, and other complaints. Other symptoms such as urinary frequency, cough, chest tightness, shortness of breath, blurred vision and night sweats were not revealed. The patient had regular menstrual cycles and no history of dysmenorrhea. Her menstrual flow was moderate, dark red and without clots. She had one daughter, and no history of cigarette smoking and alcohol abuse. The patient underwent a right ovary oophorocystectomy 10 years ago and a section of right Bartholin's cyst 4 months ago. The histopathological results of these two operations showed benign cysts.

On physical examination, we found a firm and unpainful mass (1.5 cm × 1 cm) on the edge of an episiotomy scar in the outside of the left vulva, and a callous nodule (0.5 cm × 0.5 cm) in the left vaginal wall. Other abnormal physical signs were not found. Baseline blood tests and angiotensin-converting enzyme (ACE) level were normal. The level of serum cancer antigen (CA)-125 was 43.1 U/mL (normal range <35U/mL). The PPD skin test was negative. Her spirometry showed an FVC of 3.46 L (110% predicted), an FEV1 of 2.94 L (108% predicted), an FEV1 /FVC of 0.85, an DLCO SB of 6.63 mmol/min /KPa (79% predicted). The ultrasound images of the vulva showed a low level echo of a nodule under the skin in the left vulva. There was no apparent abnormality seen in the Chest X-ray. The chest enhancement CT revealed reticulonodular infiltrates in bilateral lungs and mildly enlarged mediastinal lymph nodes (Fig. 1). Resections of the vulvar mass and vaginal wall nodule were performed.

Sarcoidosis: Vaginal wall and vulvar involvement

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Abstract. Sarcoidosis is a non-caseous granulomatous disease which could involve numerous organs including lungs, eyes, skin, nervous system, heart, liver. However, the genitourinary tract involvement was rarely reported in sarcoidosis. We report the case of a 45-year-old married woman who presented with 2 months history of a vulval mass as large as a soybean, and did not reveal any remarkable pulmonary signs. Biopsy results showed non-caseous granulomatous inflammation consistent with sarcoidosis in the vulvar lesion. To our knowledge, this is the first reported case of this entity in the world. Based on the related literature, we highlight the possibility of gynecologic involvement in sarcoidosis. (Sarcoidosis Vasculitides Diffuse Lung Dis 2012; 29: 151-154)

Key words: sarcoidosis, gynecologic involvement
Pathological examinations showed non-caseous granulomatous inflammation consistent with sarcoidosis in both the genital mass and the vaginal wall nodule (Fig. 2). PAS staining and acid-fast staining were negative for fungi and acid fast bacilli. Consistently, cultures for fungi and mycobacteria were also negative. A diagnosis of vaginal wall and vulval sarcoidosis is made after excluding other potential causes of granulomatous inflammation and cancer. Based on the staging result of the chest radiography (stage 0), glucocorticoid was not used to treat the patient. Presently, the patient is closely followed every 1 month.

**Discussion**

Sarcoidosis is a multisystem granulomatous disease affecting many organs, mostly the lungs. The pathological features are noncaseous necrotic granuloma, which was first described in 1899 by Hutchinson. (1) The cause of sarcoidosis remains unclear. It may be related to the environment, genetic predisposition and disturbance of immune function for CD4+ T cells.

Sarcoidosis is clinically manifested with pulmonary involvement. Other commonly affected organs include eyes, skin, nervous system, heart, and liver. The male patient with sarcoidosis may present with a painless testicular swelling when genitourin-
Sarcoidosis: vaginal wall and vulvar involvement

Table 1. Published sarcoidosis cases involving the vulva or vagina

<table>
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<tr>
<th>Cases</th>
<th>First involvement</th>
<th>Other involvements</th>
<th>Chief complaints</th>
<th>Images of lung (CXR or CT)</th>
<th>Diagnosis</th>
<th>Treatment</th>
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<tr>
<td>1</td>
<td>Vulva</td>
<td>Peri-anal skin</td>
<td>Papular rash on the vulva</td>
<td>Bilateral hilar lymphadenopathy and nodular shadowing in the lung fields</td>
<td>Vulval sarcoidosis</td>
<td>No record</td>
<td>No record</td>
<td>Tatnall, 1985 (4)</td>
</tr>
<tr>
<td>2</td>
<td>Vulva</td>
<td>Lung, facial skin</td>
<td>Pruritic and burning facial and vulvar eruption</td>
<td>Normal</td>
<td>Vulval involvement in sarcoidosis</td>
<td>Hydroxychloroquine, chloroquine and corticosteroid</td>
<td>The vulval lesions persisted, but improved slightly</td>
<td>Klein, 1998 (5)</td>
</tr>
<tr>
<td>3</td>
<td>Vulva</td>
<td>Lung</td>
<td>Persistent itchy papules in the vulva</td>
<td>Hilar adenopathy and a bilateral reticulonodular pattern, bilateral multiple rounded pulmonary nodules</td>
<td>Sarcoïdosis</td>
<td>Clobetasol propionate, flurandrenolone</td>
<td>Pulmonary and nodal disease has remained static</td>
<td>Ezugha, 2005 (6)</td>
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<tr>
<td>4</td>
<td>Vulva</td>
<td>Lung</td>
<td>Lesion associated with an scar in the left perineal area</td>
<td>Bilateral hilar lymphadenopathy</td>
<td>Vulval involvement in sarcoidosis</td>
<td>No treatment</td>
<td>Patient was still well, 12 months after operation</td>
<td>Decavalas, 2007 (7)</td>
</tr>
<tr>
<td>5</td>
<td>Vagina</td>
<td>Lung</td>
<td>Vaginal itching, irritation cough, wheezing on exertion, and night sweats</td>
<td>Bilateral hilar and mediastinal lymphadenopathy with diffuse multiple nodular opacities</td>
<td>Vaginal involvement in sarcoidosis</td>
<td>Corticosteroids and chloroquine</td>
<td>Clinical symptom improved, but still needed prednisone therapy</td>
<td>Allen, 2010 (3)</td>
</tr>
</tbody>
</table>

The mass of sarcoidosis was found in the episiotomy scar of the left vulva in this case, in line with another study showing a skin scar is often involved. (8) The laboratory examination showed elevated CA-125, a glycoprotein expressed by a variety of tissues of mesothelial origin. It was reported that patients with sarcoidosis in the peritoneum had higher levels of CA-125 than normal. (9) Sarcoidosis generally has a good prognosis. Sixty percent of patients do not require treatment and the disease may spontaneously regress in some patients. Glucocorticoid and other immunosuppressants have been used to treat sarcoidosis for many years. However, their influence on the natural history of sarcoidosis is unclear. Our case had no bothersome signs and her vital organs were not at risk, therefore aggressive treatment was not adopted for her. After 1 year’s follow-up, this patient is still in good condition.
REFERENCES