A Case of Multiple Myeloma with Extramedullary Plasmacytoma Involving the Testis and Pleura at the Time of Initial Diagnosis

INTRODUCTION

Extramedullary plasmacytoma (EMP) is defined as extra-osseous proliferation of neoplastic plasma cells in multiple myeloma (MM). A 55-year-old man presented with painless swelling of his left testis. Chest imaging showed plaque-like pleural thickening in the right hemithorax, and bone imaging revealed multiple sites of bony involvement. Left orchiectomy and pleural biopsy were performed and the histologic assessment revealed a diagnosis of EMP. Although testes and pleura are rarely involved in MM, the possibility of testicular or pleural EMP should be considered when a patient with MM presents with testicular swelling or a chest imaging abnormality.

Index terms
Plasmacytoma
Multiple Myeloma
Testis
Pleura

CASE REPORT

A 55-year-old man presented with painless swelling of his left testis. On testicular ultrasonography (US), the left testis measured $3.3 \times 6.7 \times 4.5$ cm in size and $51.9 \text{ cm}^3$ in volume, and showed heterogeneous echogenicity with multiple, ill-defined, hypoechoic foci and hypervascularity, which was suggestive of malignancy (Fig. 1). Chest radiography performed for a preoperative work-up showed increased opacity in the right lower pleural space.
Extramedullary Plasmacytoma Involving the Testis and Pleura

Fig. 1. A 55-year-old man who presented with painless swelling of the left testis. Gray-scale (A) and color Doppler (B) images of testis ultrasonography show heterogeneous echogenicity with multiple, ill-defined, hypoechoic foci and hypervascularity in the left testis.

Fig. 2. Chest CT scan shows plaque-like pleural thickening in the right lower posterior hemithorax (arrows).

Fig. 3. A 55-year-old man who underwent left orchietomy. A. Photograph of surgical specimen obtained from left orchietomy shows no definite nodular or mass-like lesion in testis. B. Photomicrograph of surgical specimen obtained from left orchietomy shows relatively uniform plasma cells with eccentrically located nuclei and the residual seminiferous tubules separated by heavy infiltrates of plasma cells (H&E, x 400). C, D. Immunohistochemical stains of surgical specimen obtained from left orchietomy show positive reaction for cytoplasmic lambda light chain in plasma cells (x 400) (C), and negative reaction for kappa light chain in plasma cells (x 400) (D).

lung zone. Chest CT scans performed for further evaluation showed plaque-like pleural thickening in the right lower posterior hemithorax, which was suggestive of pleural metastasis (Fig. 2). During the diagnostic work-up, the patient also complained of painless scalp swelling. Subsequent skull series revealed multiple, punched-out, radiolucent lesions in the skull. On brain CT scans, osteolytic bony lesions with focal scalp swelling were seen in the left parietal area, which were suggestive of bony metastasis. Whole-body bone scan showed multiple areas of uptake in the skull, right 4th, 9th, and 10th ribs, and right proximal femur, which were suggestive of multiple bony metastases. To confirm the diagnosis and to direct treatment, left orchietomy and pleural biopsy were performed. The histopathologic specimens obtained from left orchietomy and pleural biopsy revealed plasma cell infiltration of the interstitium of the testis (Fig. 3) and pleura (Fig. 4), positive immunohistochemical staining for CD79a, Bcl-2, cyclin D1, and lambda, and negative staining for CD30 (Ki-1), kappa, and Ki-67 (30%). Taken together, these findings confirmed the diagnosis of EMP (Fig. 3). Serum and urine protein electrophoresis revealed monoclonal gammopathy with a positive M-spike. Serum and urine immunofixation electrophoresis (IFEPE) demonstrated an abnormal band of restriction in the lambda light chain. Bence-Jones lambda chains were detected in the urine. On repeated serum IFEPE, monoclonal gammopathy of IgD type was detected. Serum and urine beta-2 microglobulin levels were 4.7 mg/L and 7.00 mg/L, respectively. Bone marrow (BM) aspirate showed a high number of plasma cells (44.6%), and BM biopsy specimen showed compact cellularity (100%) with diffuse plasma cells and scanty normal hematopoietic cells. A final diagnosis of IgD-type MM with EMP involving the testis and pleura was made. The patient...
achieved a very good partial response after two cycles of high-dose dexamethasone chemotherapy; and two cycles of vincristine, doxorubicin, and dexamethasone chemotherapy; however, the disease relapsed after auto-peripheral blood stem cell transplantation. After the patient received the fifth cycle of chemotherapy with bortezomib and dexamethasone, he had complete remission with negative serum and urine IFEP results.

**DISCUSSION**

According to the World Health Organization classification of plasma cell neoplasms, plasmacytoma is classified into solitary plasmacytoma of bone and EMP, which presents as a soft tissue mass without any other evidence of myeloma. Some of the researches have defined EMP as a plasma cell neoplasm of soft tissue with or without MM (3, 4, 6). EMP is defined as extra-osseous proliferation of neoplastic plasma cells, and may be primary or secondary to bone marrow involvement in MM. EMP is rare but most often occurs in the nasopharynx, upper respiratory tract, or gastrointestinal tract. The testis and pleura are unusual sites of EMP (1). Testicular EMP has been reported in 0.6–2.7% of all MM cases (2). According to a review of 958 MM cases, only 0.8% of patients were reported to have myelomatous pleural effusion (3). In a study assessing the clinical correlates and prognostic relevance of EMP in the presence of MM, 42 patients had EMP among 467 patients with MM (4). Of these 42 patients, only 9 patients had pleural or peritoneal EMP at the time of diagnosis and none of the patients had testicular EMP.

The clinical incidence of testicular plasmacytoma is reported to be 0.03% to 0.1% of all primary and secondary tumors of the testes (7). It is well known that testes are rare sites of hematologic malignancy because of the blood-testis barrier. Until 2002, 51 cases of testicular plasmacytoma were reported in the literature, either as the initial presenting symptom or not as the initial presenting symptom (8). In our case, the chief complaint of the patient was testicular enlargement, which proved to be testicular EMP involvement of IgD-type MM. Although our patient had a rare IgD-type MM, the US findings were similar to those of other MM types; an enlarged testicle with poorly circumscribed intra-testicular hypoechoic lesions and increased vascularity (8).

Kintzer et al. (3) reported that only eleven (1%) of 958 MM cases had intrathoracic EMP without bone involvement, and only 0.8% of the patients had myelomatous pleural effusion although the evidence of pleural EMP was unclear among these patients. Most of the pleural manifestations of MM have been reported as myelomatous pleural effusion (3, 9). In a few case reports, the CT findings of pleural EMP included focal pleural thickening or diffuse confluent pleural nodularity mimicking mesothelioma (9). In our case, the patient had plaque-like pleural thickening in the right lower posterior hemithorax on the initial CT scan, which was suggestive of pleural metastasis.

In most of the patients with MM, IgG or IgA are the immunoglobulin types that are reported; whereas in our case, the immunoglobulin type was IgD. IgD MM is a rare form of MM, which comprises 2.1% of all myelomas, is predominant in males, and occurs more frequently in younger people (10). The main characteristics of IgD MM include uncertain appearance of the M-component, dominance of the lambda light chain over the kappa chain in serum electrophoresis, and frequent presence of Bence-Jones proteins in urine. IgD MM has been known to be associated with a higher frequency of renal failure than the other types of myelomas. In our case, the patient did not have any evidence of renal failure although results of all serum and urine studies were consistent with IgD MM.

There is only one case report in the literature of MM synchronously involving the testis and pleura according to Pubmed and Medline databases. Giuliani et al. (5) reported a case of IgG MM presenting with testicular swelling and myelomatous effusion after 5 cycles of chemotherapy. In their case, the pleural involvement in MM was proved by myelomatous pleural effusion without a soft tissue mass; therefore, it was not definitive for EMP.

**Fig. 4.** Photomicrograph of biopsy specimen obtained from pleural excision shows plasma cells infiltrating the pleura (H&E, × 400).
However, in our case, the patient presented with testicular swelling and plaque-like pleural thickening at the time of initial diagnosis of MM. These soft tissues were pathologically confirmed as EMP of the testis and pleura. EMP has been reported to occur before the diagnosis of MM in only 0.4% of patients with MM (6). In addition, among patients with IgD MM, there are a few reports of patients with extraosseous disease (11).

In conclusion, we report a case of IgD-type MM with testicular and pleural EMP at the time of initial diagnosis. We think that this is the first case report of IgD MM with synchronous involvement of the testis and pleura. Although the testis and pleura are rarely involved in MM, the possibility of testicular or pleural EMP should be considered when a patient with MM presents with testicular swelling or a chest imaging abnormality.

REFERENCES

초기 진단시 고환과 흉막의 골수외 형질 세포종을 동반한 다발성 골수종의 증례 보고

신 맥 · 정세민 · 문영태 · 이태진

골수외 형질 세포종은 다발성 골수종에서 골 밖에서 발생하는 종양성 형질세포 증식이다. 55세 남환자 좌측 고환의 무통성 부종을 주소로 내원하였다. 흉부 영상에서 우측 흉곽 내에 반상의 흉막비후가 보였고 골 영상에서 다발성 골 침범이 동반되었다. 좌측 고환절제술과 흉막 생검에서 골수외 형질 세포종으로 진단되었다. 비록 다발성 골수종이 고환과 흉막을 드물게 침범하더라도, 다발성 골수종 환자에서 고환 부종 또는 흉부 이상을 동반할 때 고환 또는 흉막의 골수외 형질 세포종의 가능성이 고려되어야 한다.

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