Unilateral Diaphragm Paralysis Possibly Due to Cervical Spine Involvement in Multiple Myeloma

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Introduction

Multiple myeloma (MM) is a neoplastic disorder of plasma cells which accounts for 10% of all hematological cancers [1]. Vertebral involvement is quite common and MM patients are frequently admitted to the hospital with initial complaints of back pain. If MM is not included in the differential diagnosis, the underlying etiology for the back pain may be overlooked and the patient might undergo unnecessary interventions. In this report, we present an MM patient who had been operated upon for back pain and vertebral compressions in another hospital without a specific diagnosis of MM.

Case Report

A 52-year-old female was admitted with complaints of dyspnea, back pain and widespread bone pain. Previously, she had been given alendronate and calcium supplements along with a low back exercise program. Subsequently, she was examined by a neurosurgeon who suggested surgery for the diagnosis of 'lumbar disk herniation'. She refused the operation and later was evaluated by several orthopedic surgeons. Eventually, she underwent surgery for the vertebral compressions using kyphoplasty. Unfortunately, no pathological diagnoses were available from surgical specimens. Her back pain did not subside after these treatments. A chest physician examined her for complaints of dyspnea. Labora-
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In this patient, laboratory studies disclosed anemia and a decreased albumin/globulin ratio. These results together with her history of compression fractures suggested the need for further evaluation. The physical examination was consistent with malaise, palpitation, dyspnea, tachypnea and absence of pulmonary sounds in the left lower pulmonary lung fields.

The laboratory analysis revealed the following: hemoglobin, 11.1 g/dl (normal 12–16); leukocyte count, 6,500/µl (4,000–10,000); platelets, 250,000/µl (150,000–400,000); erythrocyte sedimentation rate, 74 mm/h (0–20.0); creatinine, 1.0 mg/dl; blood urea nitrogen, 30 mg/dl (4.6–23); total protein, 10.2 g/dl (6.0–7.8); albumin, 3.2 g/dl (3.2–4.8); globulin, 7.00 g/dl (1.5–4.6); α2-microglobulin, 4.34 (0.9–2.0). Serum protein electrophoresis yielded: γ-globulin, 48.8% (oligoclonal peak); albumin, 30.7%; α1, 5.6%; α2, 9.2%; β-globulin, 5.7%. Using immune-fixed electrophoresis, oligoclonal light chains were observed in urinalysis. A chest X-ray depicted a left-sided elevation of the diaphragm (fig. 1). Computed tomography of the thorax revealed an elevation of the diaphragm on the left side with decreased lung volume. During fluoroscopy the left diaphragm was elevated and motionless throughout the respiration cycles. The tidal volume was 460 cm³. Skeletal survey revealed multiple lytic lesions in the skull, ribs, pelvis, humerus and both femurs with generalized osteopenia. The cervicothoracolumbar magnetic resonance imaging demonstrated diffuse vertebral involvement and multiple compressions, predominantly of the cervical vertebrae (fig. 2a, b). Bone marrow biopsy revealed abnormal plasma cell infiltration. As a result, the patient was diagnosed as having MM with generalized bony involvement and left-sided diaphragmatic paralysis due to cervical cord and/or nerve compression.

The patient received two cycles of a chemotherapeutic regimen consisting of vincristine-Adriamycin-dexamethasone. The bisphosphonate zoledronic acid, 4 mg i.v. every 28 days, was also initiated. During the follow-up, the back pain resolved and the complaint of dyspnea decreased although the paralysis persisted.

**Fig. 1.** The chest X-ray of the patient demonstrating the unilateral elevation of the diaphragm on the left side.

**Fig. 2.** a Magnetic resonance imaging T1-weighted sagittal view of the cervical vertebrae depicting the extensive lytic lesions and several compressions. b T1-weighted sagittal view of the thoracic vertebrae depicting the extensive lytic lesions and several compressions.
Discussion

MM is a hematological malignancy with abnormal monoclonal plasma cell proliferation, skeletal destruction, renal failure, anemia and hypercalcemia. The most common symptoms on presentation are fatigue, bone pain and recurrent infections. Patients who are eligible for autologous stem cell transplantation are first treated for pain and recurrent infections. Patients who are eligible are most often treated with thalidomide and new drugs such as bortezomib and CC-5013. Moreover, bisphosphonates, namely zoledronic acid and pamidronate, have a widely recognized antiosteoclastic activity that decreases skeletal destruction by 50% in advanced MM.

Abnormal plasma cells proliferate in the bone marrow and then invade the bone causing lytic lesions. The lytic lesions, osteoporosis and subsequent pathological fractures are present in 75% of patients. They are often responsible for the complaints of various types of bone pain which exist in two thirds of the patients at the time of diagnosis. Involvement of the vertebrae may result in compression fractures and vertebral collapse. Besides causing back pain, these complications may frequently cause compression of the spinal cord or the spinal nerves. Otherwise, peripheral neuropathies are an uncommon finding in MM. If present, these neuropathies result from immunoglobulins reacting against myelin-associated glycoproteins and peripheral nerve antigens or to amyloid deposition. Other neurological symptoms of MM include headache, blurring of vision, precomatose, vertigo, ataxia, seizures and paraplegia. To the best of our knowledge, unilateral phrenic nerve involvement in MM has not been reported in the literature. In this patient, the left phrenic nerve might have been injured due to the involvement of MM in the cervical vertebrae, resulting in compression of the cerebral roots forming the left phrenic nerve. Unfortunately, phrenic nerve conduction studies, diaphragmatic electromyography and imaging to visualize structural damage to the ventral roots comprising the phrenic nerves were not done. However, since our patient had no accompanying neurological complaints, an infiltrative cause or immunological reaction leading to polyneuropathy was considered unlikely. Other causes of unilateral diaphragmatic paralysis include tumors, more commonly bronchogenic lung cancer, postsurgical complications, neurological disorders (myelitis, encephalitis, poliomyelitis) and herpes zoster, but these were ruled out in our patient.

A plication operation for the diaphragm was not recommended for our patient since her dyspnea resolved after chemotherapy and her tidal volume was 460 cm³.

Conclusion

This case shows that phrenic nerve paralysis from cervical vertebral compression may be an unusual complication of MM. Additionally, it is important to include MM in the differential diagnosis of back or generalized bone pain, which will facilitate the proper diagnosis and prompt treatment of this disease.

References