Primary Neuroendocrine Tumor of the Prostate with Bone Metastasis

Kemik Metastazı Yapmış Prostatın Primer Nöroendokrin Tümörü

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Özet

Anahtar Kelimeler
Prostat; Nöroendokrin; Tümör; Kemik; Metastaz

Abstract
Primary neuroendocrine tumors of the prostate are very rare and their biologic behaviour is not yet well described. A case of primary, prostatic, neuroendocrine tumor with bone metastasis is described. A 50-year-old man presented with a 6 months history of frequent urination, difficulty in urination and failure to empty the bladder. Transurethral prostate resection was performed and the pathological examination of the prostate tissue revealed a poorly-differentiated neuroendocrine carcinoma. Meanwhile, cervical magnetic resonance imaging that was planned due to the presence of neck pain revealed a mass lesion in the right paravertebral soft tissue and a pathological compression fracture of the C6 vertebral body. The patient underwent an operation in which C6 corpectomy, placement of homogenous bone graft and obtaining a biopsy specimen were performed. The pathological assessment of the biopsy revealed metastasis of a poorly-differentiated neuroendocrine carcinoma confirming the bone metastasis of the primary neuroendocrine tumor of the prostate.

Keywords
Prostat; Nöroendokrin; Tümör; Kemik; Metastaz
Introduction
Adenocarcinomas constitute approximately 95% of all malignant tumors of the prostate, and although rare, neuroendocrine tumors of the prostate, such as carcinoid tumors or small-cell carcinomas may also be encountered [1-4]. Herein, a patient diagnosed with a primary neuroendocrine tumor of the prostate in our clinic in whom evidence of bone metastasis was subsequently detected is presented.

Case Report
A 50-year-old male patient was admitted to our clinic with the complaints of frequent urination, difficulty in urination and failure to empty the bladder persisting for 6 months. His medical history revealed an acute urinary retention 3 months ago, requiring treatment with a urinary catheter and an alpha-adrenergic blocker. The patient, for whom a surgical intervention was planned, underwent a rectal examination, which revealed a soft, smooth prostate of grade I enlargement. The prostate-specific antigen (PSA) level was 1.4 ng/mL and creatinine was 1.1 mg/dL. Urinalysis revealed no abnormal findings. Prostatic obstruction was detected on cystoscopy, and a transurethral prostate resection was performed. Meanwhile, a cervical magnetic resonance imaging that was planned due to the presence of neck pain revealed a mass lesion in the right paravertebral soft tissue and a pathological compression fracture of the C6 vertebral body (Figure 1). Subsequently, the patient underwent an operation in which C6 corpectomy, placement of homogenous bone graft and obtaining a biopsy specimen were performed. The pathological examination of the prostate tissue revealed a poorly-differentiated neuroendocrine carcinoma. The immunohistochemical analysis of the tumor cells revealed positivity for CD56, synaptophysin and pancytokeratin, but negativity for PSA, CD45 and CD99 (Figure 2). The pathological assessment of the materials obtained from the C6 corpectomy and biopsy revealed metastasis of a poorly-differentiated neuroendocrine carcinoma. The immunohistochemical analysis revealed focal, weak positivity for chromogranin, diffuse, strong positivity for CD56 and synaptophysin, and negativity for CD45 (Figure 3). The patient was referred to the medical oncology clinic due to the metastatic primary neuroendocrine tumor of the prostate.

Discussion
Primary neuroendocrine tumor of the prostate is a rarely seen tumor that originates from cells responsible for amine precursor uptake and decarboxylation [4]. These tumors most often metastasize to the lymph nodes, to the liver, lungs and bones [5]. Numerous cases have been described in the literature, most of which involve cases over 60 years of age. In the presentation of a case of pediatric age, a primary neuroendocrine tumor of the prostate in conjunction with multiple endocrine neoplasia IIb was described [6]. Furthermore, the incidence of neuroendocrine tumors is increased in individuals with Crohn’s disease [7]. Moreover, neuroendocrine differentiation can be observed in 10%-100% of prostatic adenocarcinomas. Neuroendocrine cells are present in the prostate tissue, and display a widespread distribution in the prostate tissue at birth [1,8,9]. Prostate tumors that show neuroendocrine differentiation are considered to be subtypes of prostatic adenocarcinomas (9). Some authors claim that tumors with neuroendocrine differentiation behave more aggressively and hence, it is necessary to differentiate conventional adenocarcinomas with neuroendocrine differentiation from pure neuroendocrine carcinomas [10]. The present case is a pure and metastatic neuroendocrine tumor of the prostate. The fundamental treatment of pure primary neuroendocrine tumors of the prostate is radical prostatectomy. However, as in the present case, other treatment modalities are recommended in the presence of metastasis. While chemotherapeutic agents such as cisplatin and etoposide can be recommended in anaplastic variants, a good prognosis can be observed in well-differentiated tumors with no additional treatment [11]. Besides, radiotherapy can be applied for local tumor control and analgesia in the treatment of tumors of large volume, brain metastases, potential pathological fractures, and for bladder outlet obstructions. Due to the limited number of cases in the literature, our knowledge regarding the prognosis of advanced stage disease is thus far insufficient. As seen in the present case, the possibility of the presence of a neuroendocrine tumor of the prostate should also be kept in mind in patients presenting with lower urinary tract complaints who have normal PSA levels.

Competing interests
The authors declare that they have no competing interests.

References

Figure 1. Magnetic resonance imaging showing pathological compression fracture of the C6 vertebral body and metastatic mass lesion (arrow) in the right paravertebral soft tissue

Figure 2. Immunohistochemical staining. (A) Neoplastic cells showing positivity for synaptophysin, (B) and for CD56

Figure 3. Islets of neoplastic cells (arrow) between the bone spicules