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Ectopic Cushing's syndrome in a patient with metastatic Merkel cell carcinoma: A case report

Avraham Ishay, Elia Touma, Olga Vornicova, Roni Dodik-Gad, Tal Goldman, Naiel Bisharat

BACKGROUND
Ectopic Cushing syndrome (ECS) is a rare condition commonly associated with neuroendocrine tumors (NET), mainly bronchial carcinoids. The association of paraneoplastic syndrome with Merkel cell carcinoma (MCC) is limited to individual case reports.

CASE SUMMARY
In this article we report an unusual and striking presentation of ECS in a patient with known metastatic MCC. An elderly patient presented with new onset severe hypertension, hyperglycemia and hypokalemia, muscle wasting, and peripheral edema. A diagnosis of adrenocorticotropic hormone dependent, non-pituitary, Cushing syndrome was established. Medical therapy inhibiting adrenal function was promptly started but unfortunately the patient survived only a few days after diagnosis.

CONCLUSION
The occurrence of an aggressive form of ECS in patients with NET should be recognized as an ominous event. To our knowledge, the association of this complication in a patient with MCC had not been reported.

Key Words: Merkel cell carcinoma; Paraneoplastic syndrome; Ectopic Cushing's
syndrome; Neuroendocrine tumor; Hypercortisolism; Skin cancer; Case report

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Core Tip: Merkel cell carcinoma (MCC) is an uncommon but highly aggressive skin cancer with neuroendocrine features. Its incidence and mortality are increasing. We describe an elderly patient with a 2-year history of metastatic MCC, with no apparent cutaneous lesion at diagnosis, who presented with uncontrolled hypertension, diabetes mellitus, and hypokalemia. A diagnosis of ectopic Cushing syndrome was established. The occurrence of ectopic Cushing syndrome in patients with neuroendocrine tumor is a major cause of poor prognosis. To our knowledge, this is the first reported case of ectopic Cushing syndrome linked to the rapid progression of a metastatic MCC.

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INTRODUCTION

Merkel cell carcinoma (MCC) is an uncommon but highly aggressive skin cancer with neuroendocrine features[1]. It was first described by Toker in 1972 as "trabecular carcinoma of the skin"[2]. Evidence suggests that its incidence and mortality are increasing across the world[3]. Ectopic Cushing syndrome (ECS) is a rare condition due to ectopic production of adrenocorticotropic hormone by non-pituitary tumors. The adrenocorticotropic hormone producing neoplasms usually originate from neuroendocrine tumors (NET) and can present as benign indolent tumors or aggressive metastatic tumors with a poor prognosis[4]. Although in the past elevated adrenocorticotrophic hormone levels in plasma and tumoral tissue were demonstrated in patients with MCC[5,6], there are no reports of adrenocorticotrophic hormone producing MCC that fulfill the diagnostic criteria for ECS. We describe a patient with metastatic MCC who developed an aggressive form of ECS.

CASE PRESENTATION

Chief complaints

An 82-year-old man with a 2-year history of MCC was referred for evaluation and treatment of uncontrolled high blood pressure and new onset hyperglycemia.

History of present illness

In 2018, a cervical lymphadenopathy biopsy showed metastatic MCC with no apparent primary cutaneous lesion (Figure 1). Multiple bone metastases were demonstrated by an 18F (18-fluorodeoxyglucose) PET/CT scan (18FDG-PET/CT). No pathological uptake was seen in the lungs. The patient achieved good response to avelumab initially as disclosed by a significant reduction of uptake intensity in cervical lymph nodes and in the skeleton on a subsequent 18FDG-PET/CT scan. But thereafter, the disease progressed despite adjuvant radiotherapy and systemic therapy including etoposide, carboplatin, and topotecan. Indeed, a further 18FDG-PET/CT scan showed intensification of the uptake in bones, and new metastases in mediastinal lymph nodes and multiple cutaneous lesions. Noticeably, no disease was present in the lungs.

History of past illness

The patient’s history was significant for multiple surgical treatments for squamous cell carcinoma and basal cell carcinoma of the skin.

Personal and family history

Multiple surgical treatments for squamous cell carcinoma and basal cell carcinoma of the skin.

Physical examination

Physical examination revealed high blood pressure (198/100), and a 4 cm-sized purplish-blue tumor in his central chest (Figure 2), bilateral axillary lymphadenopathy, and bilateral lower extremities pitting
edema. The clinical phenotype was dominated by weight loss and muscle wasting.

**Laboratory examinations**

Initial blood work revealed a glucose level of 241 mg/dL with hypokalemia (2.7 nmol/L). The 24-h free urine Cortisol level was 9986 nmol/24 hr (normal values: 57.7-806.8). After high dose (8 mg) overnight
Table 1 Cortisol and adrenocorticotropic hormone (ACTH) levels after stimulation with IV corticotropin releasing hormone 100 µg

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<th>Time (mn)</th>
<th>-15</th>
<th>0</th>
<th>15</th>
<th>30</th>
<th>45</th>
<th>60</th>
<th>90</th>
<th>120</th>
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<td>Cortisol (µg/d)</td>
<td>52.22</td>
<td>48.92</td>
<td>45.48</td>
<td>56.11</td>
<td>51.44</td>
<td>52.1</td>
<td>53.08</td>
<td>56.65</td>
</tr>
<tr>
<td>ACTH (pmol/mL)</td>
<td>92.9</td>
<td>110</td>
<td>129</td>
<td>128</td>
<td>122</td>
<td>96.1</td>
<td>119</td>
<td>120</td>
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dexamethasone suppression test (HDST), the 8-am serum cortisol was 44.9 µg/dL (normal values: 4.3-22.4). Serum adrenocorticotropic hormone level was elevated: 106 pg/mL (0.0-46). Testing with intravenous corticotropin releasing hormone (CRH) administration did not affect adrenocorticotropic hormone or cortisol levels, which is typical of ECS (Table 1)

**Imaging examinations**
Magnetic resonance imaging scan of the pituitary gland was normal.

**FINAL DIAGNOSIS**
Ectopic adrenocorticotropic hormone dependent Cushing’s syndrome.

**TREATMENT**
The patient was treated with ketoconazole.

**OUTCOME AND FOLLOW-UP**
The patient’s blood pressure and glucose levels were normalized; however, his general state did not allow additional antineoplastic therapy and he died within few days.

**DISCUSSION**
We present an 82-year patient with a metastatic MCC presenting with an overwhelming form of ECS. ECS is a rare condition which accounts for about 10%-20% cases of adrenocorticotropic hormone dependent Cushing syndrome. Neuroendocrine tumors (NETs), principally bronchial carcinoids, are the most frequent causes of ECS. Less frequent causes are thymic carcinoids and pancreatic NETs [7]. Small cell lung carcinoma is a known cause of ECS, but in our patient imaging studies did not reveal any lung lesions. Recently, a case of metastatic NET of unknown origin presenting with ECS was reported [8]. Several manifestations of MCC-associated paraneoplastic syndromes have been reported [9], but ECS associated with MCC has not be described, even though a case of metastatic MCC within a cortisol-producing adrenal adenoma has been recently reported [10]. The time elapsed between the first symptoms of hypercortisolism and the diagnosis of ECS may predict the prognosis of the underlying malignancy. The shorter it is, the poorer is the prognosis. In addition to the grade of NET, the severity of cortisol excess is an independent negative prognostic factor [7]. The molecular mechanisms underlying ECS-associated malignant tumors include aberrant processing of the proopiomelanocortin (POMC) gene leading to release in the circulation of high molecular weight adrenocorticotropic hormone precursors like POMC and pro-adrenocorticotropic hormone. It is speculated that the ability of the tumor to express aberrant molecules is related to the progression of the disease [11]. If ectopic adrenocorticotropic hormone producing malignancy is diagnosed early as a localized disease, surgical removal of the primary tumor is the treatment of choice, but it is rarely achievable in patients with aggressive neoplasms. In this ECS group, a prompt control of hypercortisolism should be attempted by medical treatment or alternatively by adrenalectomy [4].

**CONCLUSION**
MCC and neuroendocrine ECS are both rare conditions. The occurrence of ECS in patients with metastatic NETs is a major cause of poor prognosis. The suspicion of Cushing syndrome should receive adequate attention and prompt evaluation to confirm the diagnosis and initiate rapidly the treatment to
attain a more favorable prognosis.

**FOOTNOTES**

**Author contributions:** Ishay A and Touma E were the patient's physicians, wrote the manuscript, and reviewed the literature; Vornicova O, Doduk Gad, and Bisharat N were involved in the patient treatment and contributed to manuscript drafting; Goldman T was the pathologist and prepared and interpreted the pathology images.

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