Near missed diagnosis of Merkel Cell Carcinoma in a young immunocompetent woman with a recurrent left-arm mass: A case report

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INTRODUCTION

Merkel Cell Carcinoma (MCC) is a rare, life-threatening tumor with neuroendocrine features. Due to its extremely low occurrence worldwide, epidemiological data on the disease is limited. Commonly recognized risk factors include fair skin, history of skin cancer, old age, chronic immunosuppression, chronic ultraviolet (UV) light exposure, and Merkel cell polyomavirus (MCPyV) infection. Amongst these factors, MCPyV and UV exposure play a fundamental role with synergistic effects in the pathophysiology of MCC. Despite being a part of normal skin flora in most individuals, MCPyV DNA can clonally combine with the genome of neoplastic cell precursors at the initial phases of carcinogenesis. Concurrently, UV exposure elicits antigen-presenting dendritic cells to produce inflammatory cytokines, leading to local immunosuppression and creating an ideal environment for tumoral growth. Moreover, immune hypersensitivity from UV exposure facilitates the viral tumorigenic process.

Merkel cells cannot be derived from the MCC because they lack the ability to proliferate. Merkel cell precursors (perhaps generated from epidermal stem cells or hair follicle stem cells) and pre- and pro-B cells appear to have histopathology, genetics, and molecular characteristics with malignant cells instead.

MCC often presents as a single, asymptomatic erythematous or violaceous nodule, often mistaken for cysts or abscesses. It usually originates from the head or neck and generally spares the extremities. Immunohistochemical (IHC) staining is required to validate histopathological findings of small round cells that infiltrate cutaneous or subcutaneous area.

Here, we present a 32-year-old Iranian (non-Caucasian) immunocompetent female with a small nodule on her left arm at the disease onset finally diagnosed as MCC. In this report, we aim to emphasize the significance of early diagnosis and management of this cancer and highlight the complications that a late diagnosis would entail for these patients.

CASE REPORT

A 32-year-old female with an unremarkable medical history presented to the clinic with a small, non-tender, and erythematous nodule on the dorsolateral aspect of her left arm, which initially appeared three months before. Since then, the nodule had slowly darkened and grown to 1 cm × 1 cm in size. Her history and
physical exam were insignificant except for multiple warts on the dorsal aspect of the right hand. (Figure 1). Initially, the patient was suspected of having an infectious cyst and was treated with 10-days of antibiotics. However, she was unresponsive to antibiotics and was evaluated by a surgeon, who diagnosed the lesion as an abscess and surgically removed it.

After two weeks, a rapidly growing mass measuring 4 cm × 4 cm originated from the incised area. The mass was surgically excised again but reappeared within one week, measuring 5 cm × 5 cm, after which the patient was referred to our hospital for further evaluation. The magnetic resonance imaging (MRI) of the left arm with and without contrast showed multiple enhancing lesions at the subdermal region of the posterolateral aspect of the left arm with multiple enlarged axillary lymph nodes, the largest of which measured approximately 17 mm, suggestive of metastasis. The tumor was radically excised with negative surgical margins. Microscopic examination of the lesion demonstrated a neoplastic round cell tumor with prominent foci of necrosis (Figure 2). IHC staining revealed neoplastic cells with positive expression for CD99, Ck20, and NSE (Figure 2), but no LCA, Vimentin, CD3, CD20, and HMB45 expression consistent with MCC (Figure 3). Also, The MCPyV PCR real-time of the lesion was positive.

High-resolution computed tomography (HRCT) Scan of the chest and abdominopelvic area with and without contrast showed lymphadenopathy 39 mm × 25 mm in the left axillary fossa along with a calcified lymph node (6mm) in the abdominal cavity beyond the left abdominal muscle. She was diagnosed with stage III MCC and subsequently treated with adjuvant radiation to her left arm and axilla with 45.0 Grays (Gy) of radiation in 35 fractions, followed by six cycles of chemotherapy with etoposide and cisplatin. The patient’s post-treatment PET/CT scan showed no evidence of neoplastic disease and supported remission. However, high-dose chemotherapy caused frequent myalgias and weakened immune system. Unfortunately, she developed febrile neutropenia and expired due to septic shock one month after the last chemotherapy course.

DISCUSSION

MCC is a rare, aggressive malignancy with an estimated annual incidence rate of 0.01 to 0.13 per 100,000 people globally. It predominantly affects the elderly (probably due to chronic UV radiation from the sun exposure), Caucasian males, and chronically immunocompromised patients, especially recipients of organ transplants, those with lymphoproliferative disorders, and untreated HIV infection (1). Specifically, in the United States, more than 8 in 10 individuals diagnosed with MCC are older than 70 years old, and more than 95% are Caucasian. Furthermore, males are twice more likely to be diagnosed with MCC than females (10). When MCC occurs in a younger population, it often involves children and is extremely rare in middle-aged adults (11).

Because of its low incidence, particularly in non-Caucasian populations, asymptomatic nature, and indistinguishable clinical presentation, MCC has a high rate of misdiagnosis (12). A retrospective analysis of 195 patients diagnosed with MCC found that less than 1% were suspected of having MCC on clinical evaluation, leading to a median delay of more than three months from the initial appearance of the nodule to biopsy sampling (13). Similarly, our patient was initially suspected of having an infectious lesion rather than MCC, which significantly delayed her initial presentation until her diagnosis. This case was further complicated by the absence of any risk factors in our patient. The most commonly recognized modifiable risk factors include MCPyV and long-term UV radiation exposure, with more than half of primary MCC lesions originating from the head and neck region (14). Despite the presence of MCPyV in her arm lesion and multiple warts on the dorsal aspect of both hands, she had none of the previously mentioned risk factors.

The clinical presentation of MCC is often variable and nonspecific. MCC often presents as a tender, erythematous red to the violet-colored lesion on sun-exposed skin regions, most commonly the head and neck, but less commonly on the trunk or the extremities (15). The only distinguishing characteristic of MCC is its rapid growth rate. Otherwise, the lesion may or may not have central ulceration and may present with superimposing infection, resulting in its misdiagnosis as an abscess (16, 17). Tender or non-tender painless
local or distant lymphadenopathy may also be present in case of lymph-node metastasis or superimposed infection (18).

Barreira et al. reported a 70-year-old immunocompromised woman with painless inguinal lymphadenopathy. Further evaluations revealed a pink plaque in the left knee whose histopathology confirmed MCC with lymph node metastasis. Like our patient, she died due to a high tumor stage and metastasis at the disease onset. However, palliative treatment was indicated for this patient since she was dealing with underlying medical problems such as nephrectomy due to renal tuberculosis and non-Hodgkin’s lymphoma (19). Similarly, Agut-Busquet E et al. reported a young white woman with a well-defined subcutaneous mass measuring 3 cm × 2.5 cm in size located in the dorsal aspect of the left arm. Nevertheless, MCC was diagnosed before the tumor spread, resulting in more effective treatment and complete remission one year after the diagnosis (20).

Diagnostic imaging, including regional lymph nodes ultrasonography, CT scan, MRI, and PET-CT scan, are often used for clinical staging and monitoring a patient’s prognosis. Early clinical detection is essential, and the possibility of MCC should be considered in patients with rapidly evolving skin lesions unresponsive to antibiotic therapy. However, neither clinical evaluation nor imaging can accurately diagnose MCC, with histopathologic evaluation and IHC studies the gold standard diagnostic approach (18). Histopathology generally displays small, uniformly rounded blue neoplastic cells with scanty cytoplasm. Even larger pleomorphic cells with increased proliferation rate, broad tissue infiltration, and lymphatic involvement may be detected. MCC-specific IHC markers should confirm the diagnosis since they distinguish this cancer from other small round cell tumors. The malignant cells show positive immunoreactivity for CK20, CK8, CK18, CK19, synaptophysin, HIP1, P36, TTF1, ASH1, S100B, and CK7, while Vimentin does not stain in the IHC of MCC (21).

Clinical manifestations of patients easily distinguish between basal cell carcinoma (BCC) and metastatic cell carcinoma (MCC). In pathology, BCC neoplastic cells display size variability and stretched nuclei with marked peripheral palisading. Unlike BCC, local lymph node metastasis and intradermal spreading are characteristic of MCC. Nevertheless, atypical cases of these two malignancies share similarities, including the presence of mucin or amyloid in the stroma and peripheral slits located in the tumor borders. Therefore, IHC plays a significant role in differentiating these challenging samples. In contrast to BCC, MCC stains with CK20 and epithelial membrane antigen (22).

Small cell melanoma is a subtype of cutaneous melanoma that displays the intraepidermal pagetoid spread in which round or atypical dendritic melanocytes gather in nests. Despite expressing S100, the presence of keratins and NSE differentiates this skin cancer from MCC (23). Although lymphoma a presents quick indistinct inflammation with prominent small cells in histology, its hematoxylinoid markers, such as PAX5, TdT, and immunoglobulins, are not detected in IHC staining of MCC (24). Also, lymphoma lacks most IHC markers of MCC, such as CD45, CD3, and CD20 (25). Also, MCC and primary cutaneous Ewing sarcoma share similarities. Small tumor cells that may be positive for keratin, CD99, FLI-1, and NSE may be seen in both types of tumors. CK20 and dot-like keratin are not found in Ewing Sarcoma while EWSR1 translocation defect is specifically detected in this malignancy (26).

Imaging techniques, including ultrasonography of regional lymph nodes, CT scan, MRI, and PET-CT scan associated with sentinel lymph node biopsy, are essential for clinical staging, prognosis, and patient follow-up. The mainstay treatment of MCC is radical surgical excision accompanied by wide-field adjuvant radiotherapy in patients with lymph node invasion (7). Chemotherapy indicated for systemic eradication of neoplastic cells often fails to restrain tumor invasion and acts as palliative care (27). Furthermore, retrospective analyses showing inconsistent results on the effects of post-operative chemoradiation on patient survival outcomes. Likewise, immune-check-point inhibitors against pathways involved in pathogenesis is reserved for advanced-stage cases unresponsive to chemotherapy (28).

In summary, we present a case of MCC with unspecific skin involvement who was misdiagnosed at first and then underwent multiple complicated surgeries. This report focused on the adverse effects of mismanagement in MCC that led to its spread and made all the therapeutic options ineffective. It underlined the consequences
of delayed diagnosis in aggressive skin tumors, as higher stages are associated with dismal prognosis despite multidisciplinary approach and patient immunocompetency. Therefore, in rapidly growing and recurrent cutaneous lesions, prompt histopathologic assessment is required to improve the patient’s overall survival and minimize side effects.

AUTHOR CONTRIBUTIONS
Fateme Salemi: Conceptualization, Methodology, Investigation, Writing- Original Draft, Writing - Review & Editing, Visualization; Seyed Mohammad Reza Mortazavizadeh: Conceptualization, Methodology, Investigation, Data curation; Project administration; Resources; Shokouh Taghipour Zahir: Conceptualization, Methodology, Resources, Writing - Review & Editing, Supervision; Soroush Shahrokh: Review & Editing.

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CONFLICT OF INTEREST STATEMENT
The authors declare no conflict of interest.

DATA AVAILABILITY STATEMENT
The case report data is not publicly available, but it could be available from the corresponding author with a reasonable request.

CONSENT
Written informed consent was obtained from the patient for publication of this case report and any accompanying images.

REFERENCES


Figures

Figure 1. multiple warts on dorsal aspect of the patient’s right hand.

Figure 2. Hematoxylin and Eosin (H&E) staining of the cutaneous lesion showing tumoral cells (10x) (a), (40x) (b).
Figure 3. IHC staining, the tumoral cells are positive for synaptophysin, (40x) (a), focally positive for S-100 (40x) (b), positive for AE1/AE3 (40x) (c), negative for Vimentin (10x) (d), positive for CD56 (40x) (e), positive for Ki67 (10x) (f), and negative for CD45 (40x) (g).