Irritated Pigmented Plaque on the Scalp

Venkata Anisha Guda, BS; Sara E. Chapman, MD; Justin P. Bandino, MD

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b. hidroacanthoma simplex
c. malignant melanoma
d. seborrheic keratosis
e. squamous cell carcinoma in situ

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Ms. Guda is from the UT Health San Antonio Long School of Medicine, Texas. Drs. Chapman and Bandino are from the Department of Dermatology, San Antonio Uniformed Services Health Education Consortium. The authors report no conflict of interest. The views expressed are those of the authors and do not reflect the official views or policy of the US Department of Defense. Correspondence: Venkata Anisha Guda, BS (gudav@livemail.uthscsa.edu).
doi:10.12788/cutis.0092
Melanoacanthoma (MA) is an extremely rare, benign, epidermal tumor histologically characterized by keratinocytes and large, pigmented, dendritic melanocytes. These lesions are loosely related to seborrheic keratoses, and the term was first coined by Mishima and Pinkus in 1960. It is estimated that the lesion occurs in only 5 of 500,000 individuals and tends to occur in older, light-skinned individuals. The majority are slow growing and are present on the head, neck, or upper extremities; however, similar lesions also have been reported on the oral mucosa. Melanoacanthomas range in size from 2×2 to 15×15 cm; are clinically pigmented; and present as either a papule, plaque, nodule, or horn.

Classic histologic findings of MA include papillomatosis, acanthosis, and hyperkeratosis with heavily pigmented dendritic melanocytes diffusely dispersed throughout all layers of the seborrheic keratosis-like epidermis. Other features include keratin-filled pseudocysts, Langerhans cells, reactive spindling of keratinocytes, and an inflammatory infiltrate. In our case, the classic histologic findings also were architecturally arranged in oval to round clones within the epidermis (quiz images 1 and 2). A MART-1 (melanoma antigen recognized by T cells) immunostain was obtained that highlighted the numerous but benign-appearing, dendritic melanocytes (quiz image 2 [inset]). A dual MART-1/Ki67 immunostain later was obtained and demonstrated a negligible proliferation index within the dendritic melanocytes. Therefore, the diagnosis of clonal MA was rendered. This formation of epidermal clones also is called the Borst-Jadassohn phenomenon, which rarely occurs in MAs. This subtype is important to recognize because the clonal pattern can more closely mimic malignant neoplasms such as melanoma.

Hidroacanthoma simplex is an intraepidermal variant of eccrine poroma. It is a rare entity that typically occurs in the extremities of women as a hyperkeratotic plaque. These typically clonal epidermal tumors may be heavily pigmented and rarely contain dendritic melanocytes; therefore, they may be confused with MA. However, classic histology will reveal an intraepidermal clonal proliferation of bland, monotonous, cuboidal cells with ample pink cytoplasm, as well as occasional cuticle-lined ducts (Figure 1). These ducts will highlight with carcinoembryonic antigen and epithelial membrane antigen immunostaining.

Malignant melanoma typically presents as a growing pigmented lesion and therefore can clinically mimic MA. Histologically, MA could be confused with melanoma due to the increased number of melanocytes plus the appearance of pagetoid spread resulting from the diffuse presence of melanocytes throughout the neoplasm. However, histologic assessment of melanoma should reveal cytologic atypia such as nuclear enlargement, hyperchromasia, molding, pleomorphism, and mitotic activity (Figure 2). Architectural atypia such as poor lateral circumscription of melanocytes, confluence and pagetoid spread of nondendritic atypical junctional melanocytes, production of pigment in deep dermal nests of melanocytes, and lack of maturation and dispersion of dermal melanocytes also should be seen. Unlike a melanocytic neoplasm, true melanocytic nests are not seen in MA, and the melanocytes are bland, normal-appearing but heavily pigmented, dendritic melanocytes. Electron microscopy has shown a defect in

**FIGURE 1.** Hidroacanthoma simplex. Intraepidermal clonal proliferation of bland, monotonous, cuboidal cells with ample pink cytoplasm, as well as occasional cuticle-lined ducts (H&E, original magnification ×200).

**FIGURE 2.** Melanoma. Large nests of variable sizes with cytologic atypia as well as confluent and pagetoid spread of junctional melanocytes (H&E, original magnification ×200).
the transfer of melanin from these highly dendritic melanocytes to the keratinocytes.

Similar to melanoma, seborrheic keratosis presents as a pigmented growing lesion; therefore, definitive diagnosis often is achieved via skin biopsy. Classic histologic findings include acanthotic or exophytic epidermal growth with a dome-shaped configuration containing multiple cornified hornlike cysts (Figure 3). Multiple keratin plugs and variably sized concentric keratin islands are common features. There may be varying degrees of melanin pigment deposition among the proliferating cells, and clonal formation may occur. Melanocyte-specific special stains and immunostains can be used to differentiate MA from seborrheic keratosis by highlighting numerous dendritic melanocytes diffusely spread throughout the epidermis in MA vs a normal distribution of occasional junctional melanocytes in seborrheic keratosis.

Squamous cell carcinoma in situ presents histologically with cytologically atypical keratinocytes encompassing the full thickness of the epidermis and sometimes crushing the basement membrane zone (Figure 4). There is a loss of the granular layer and overlying parakeratosis that often spares the adnexal ostial epithelium. Clonal formation can occur as well as increased pigment production. In comparison, bland keratinocytes are seen in MA.

Establishing the diagnosis of MA based on clinical features alone can be difficult. Dermoscopy can prove to be useful and typically will show a sunburst pattern with ridges and fissures. However, seborrheic keratoses and melanomas can have similar dermoscopic findings, therefore, a biopsy often is necessary to establish the diagnosis.

REFERENCES