Clinical Study
Desmoplastic Melanoma: Report of 5 Cases

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Background. The clinical presentation of desmoplastic melanoma is often challenging. We report the experience of the Melanoma Unit of Spedali Civili University Hospital in Brescia, Italy. Method. Study subjects were drawn from 1770 patients with histologically confirmed melanoma. Within this group, desmoplastic melanoma developed in 5 patients. For each diagnosed melanoma, histological characteristics, treatment, and outcomes were evaluated. Results. Of the 5 patients described in this study, 2 were males and 3 females. The average age was 62.4 years ranging from 56 to 68 years. Breslow thickness ranged from 2.1 to 12 mm with a mean thickness of 5.8 mm. Primary treatment of 5 patients included a wide local excision of their primary lesions. Conclusions. Desmoplastic melanoma is a rare neoplasm which clinically may mimic other tumours or cutaneous infiltrate of uncertain significance. The diagnosis is histopathological and radical resection is necessary.

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1. Background
Desmoplastic melanoma (DM) is a rare variant of malignant melanoma first described by Conley et al. in 1971 as an invasive melanoma composed of spindle cells surrounded by abundant collagen [1]. The clinical presentation of desmoplastic melanoma is often challenging. Usually it has an innocuous clinical appearance and it is described as an indurated discoid papule, plaque, or nodule. It most commonly arises in chronically sun-damaged skin. Rarely, DM may present on a mucosal surface. We describe our experience at Melanoma Unit of Spedali Civili University Hospital in Brescia, Italy, with patients whose melanoma was reported to be desmoplastic.

2. Material and Methods
Study subjects were drawn retrospectively from 1770 patients with histologically confirmed melanoma, including melanoma in situ, between January 1, 1984, and January 1, 2009. These patients were followed up by the Melanoma Unit of Spedali Civili University Hospital in Brescia, Italy. All the patients gave an informed consent to be entered into the database. A total of 5 patients (0.3%) who had been given a diagnosis of cutaneous desmoplastic melanoma, which was histologically confirmed, were identified (Table 1). All patients were staged with the use of the American Joint Committee on Cancer (AJCC) staging classification [2]. The patient’s sex, ethnic origin, age, and Fitzpatrick skin types at diagnosis were recorded. For each diagnosed melanoma, histologic classification, anatomic location, tumor thickness (Breslow method), Clark’s anatomical level of invasion, ulceration, regression, neurotropism, lymphatic invasion and number of mitoses per square millimetre were evaluated. Melanomas were classified as being located in the head and neck, trunk (including chest, back, abdomen, and buttock), upper extremity (including arm, forearm, and hand), and lower extremity (including thigh, leg, and foot). Treatment and outcomes were also examined.

3. Results
In the period between January 1, 1984, and January 1, 2009 5 patients with cutaneous desmoplastic melanoma were
identified. The incidence of desmoplastic melanoma cases in our population was 0.3% (5/1770). Of the 5 patients evaluated, 2 were males and 3 females. All patients were Caucasians. The average age was 62.4 years ranging from 56 to 68 years. Of the 5 documented patients, percentages of Fitzpatrick skin types were as follows: II 60% (3 pz) and III 40% (2 pz). Primary melanomas were found in different body sites: 1 (20%) on the trunk, 2 (40%) on the upper extremity, and 2 (40%) on the lower extremity. None of our 5 cases had a documented family history of melanoma. Breslow thickness ranged from 2.1 to 12 mm with a mean thickness of 5.8 mm. One patient had a lesion with neurotropism (case 5). Primary treatment of 5 patients included wide and deep local excision of their primary lesions. 4 out of 5 patients underwent sentinel lymph node biopsy (SLNB) and no patient was found to have a positive sentinel node. Follow-up ranged from 2 to 113 months, with an average of 49 months, and during follow-up period any patient had disease progression or developed local recurrence.

4. Conclusions

The incidence of desmoplastic melanoma cases in our population was 0.3% (5/1770). Of the 5 patients evaluated, 2 (40%) were males and 3 (60%) female. This female predominance in sex distribution is not in accordance with previous case series [3]. The median age at diagnosis of DM is approximately 10 years higher than the one for conventional melanoma [4]. In our case the average age was 62.4 years ranging from 56 to 68 years. Head and neck is the preferred site for DM; in the series from the Massachusetts General Hospital, 75% of the tumors occurred in this anatomic site [5]. None of our 5 cases had a primary melanoma in the head and neck. The clinical presentation of desmoplastic melanoma is often challenging. Usually it has an innocuous clinical appearance and it is described as an indurated discoid papule, plaque, or nodule. Pigmentation is frequently absent (Figure 1), although a lentigo or lentigo maligna-like discoloration adjacent to the nodule is not uncommon [6]. The diagnoses that are suspected clinically range from benign (scar, dermatofibroma, melanocytic nevus) to malignant (basal cell or squamous cell carcinoma, sarcoma, or amelanotic melanoma) lesions. In a series of 113 cases of desmoplastic melanoma melanoma was the initial clinical diagnosis in only 27% of the cases [3].

There are no dermoscopic criteria for this rare neoplasm probably because dermoscopic examination is not routinely performed before excising a lesion that is often clinically not considered as melanocytic. Debarbieux et al. reported the dermoscopic features for six cases of desmoplastic melanoma. In this short series only half of the cases exhibited one classical feature of a melanocytic lesion while other cases were recognized on the basis of the presence of figures of regression (all six) such as white scar-like and "peppering", multiple (>4) colours (five out of six), and melanoma-related vascular patterns (five out of six) such as linear-irregular vessels and milky-red areas [7]. The diagnosis of DM is histopathological; the histology of classic DM (Figures 2 and 3) is defined as a dermal-based, paucicellular proliferation of atypical spindle cells in a sclerotic or neuromatous stroma with evidence of melanocytic differentiation [8]. There is histological subclassification of DM into two variants: pure desmoplastic melanoma (pDM) when desmoplasia is prominent throughout the tumor and mixed desmoplastic melanoma (mDM) when desmoplasia constitutes only a part of an otherwise nondesmoplastic invasive melanoma.

The majority of DMs are larger than 1 mm in thickness at the time of diagnosis, and many tumors measure more than 4 mm. This probably results from the difficulties in clinical
also be quite different. The diagnosis is histopathological and radical resection is necessary. Current controversies regarding locoregional treatment strategies warrant further investigation.

References
