

## CONFLICTS OF INTEREST

The authors have nothing to disclose.

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# Cutaneous Plasmacytosis Showing a Neuronal Involvement in a 35-Year-Old Female

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Dear Editor:

Cutaneous plasmacytosis (CP) is a rare skin disorder characterized by multiple reddish-brown patches and nodules that mainly occur on the trunk in adults<sup>1</sup>. Histologically, there is a dermal infiltrate of mature plasma cells. We re-

port a case of 35-year-old female who presented with multiple red to brownish macules and patches on the left side of the trunk with a neuronal distribution (Fig. 1). The lesion appeared 4 years ago without any symptoms of pain or itching. A histopathological examination revealed a dense dermal patch-like cellular infiltration surrounding the follicles and the nerves (Fig. 2). The infiltrate was composed of lymphocytes, histiocytes, and plasma cells. Immunohistochemical staining was positive for CD138, CD3, and CD20. Serum protein electrophoresis and immunoelectrophoresis revealed polyclonal hypergammaglobulinemia. Based on these findings, a diagnosis of CP was made. The patient underwent treatment with systemic corticosteroids and narrowband ultraviolet B for 3 months till date. However, the response to the treatment has been minimal. In our review of the literature, skin manifestations of CP were considerably uniform among patients, with multiple reddish to brownish colored scaly patches, nodules, and

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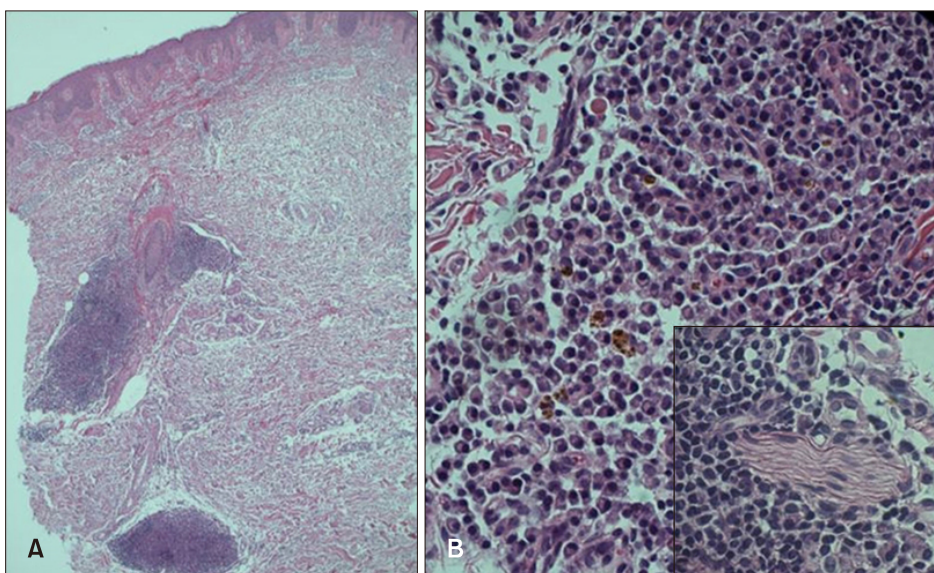
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**Fig. 1.** Multiple red to brownish maculopatches in the left side of the trunk showing a neuronal dermatosis. We received the patient's consent form about publishing all photographic materials.



**Fig. 2.** (A) Lymphoid follicle-like nodular cell infiltrates in the middle and deep dermis intermingled with sparse interstitial cell infiltrates (H&E stain,  $\times 40$ ) (B) Nodules are composed of plasma cells without atypia. Plasma cells are surrounding the neuron (H&E stain,  $\times 400$ ; inlet,  $\times 400$ ).

infiltrated plaques. Moreover, they were usually persistent, asymptomatic, or mildly pruritic and were mainly distributed on the chest or back in a symmetrical or "Christmas tree" like pattern, which may require precise differential diagnosis including pityriasis rosea and small patch parapsoriasis<sup>1,2</sup>. Several other cases with facial or scalp involvement demonstrated a typical trunk involvement pattern as well<sup>2</sup>. Interestingly, in our patient, the distribution of the lesion showed an atypical pattern, affecting only the left side of the body and showing neuronal pattern dermatoses.

The histopathologic findings characteristically show dermal nodular and perivascular/periadnexal cell infiltrations with predominance of plasma cells admixed with variable numbers of lymphocytes and histiocytes<sup>3</sup>. Recently, Honda et al.<sup>3</sup> reported 6 cases of CP all showing the perineural

and intraneural distribution of plasma cells. Similarly, in our patient's punch biopsy specimen, we observed perineural plasma cell infiltration. In this case, we hypothesize that the numerous plasma cells aggregated around the nerve fiber may have a correlation with the unilaterally neuronal pattern of the skin lesions. Furthermore, the typical symmetric pattern of CP on the chest or back may be a phenotypic expression reflecting its close relationship to the spinal nerve tract distribution.

Protein electrophoresis and immunoelectrophoresis generally show polyclonal increase in the gamma globulin fraction in major portion (84%) of the CP patients<sup>2</sup>. However, the direct relationship between polyclonality and etiology of CP is unclear. If monoclonality is observed, it is necessary to exclude the possibility of marginal zone B cell lymphoma. Polymerase chain reaction for

immunoglobulin heavy chain gene rearrangement and B-cell lymphoma 2 staining would be helpful to make the diagnosis clear<sup>4</sup>.

In summary, a diagnosis of CP should be considered when dermatologists encounter cases of asymptomatic neuronal pigment dermatosis. We speculate that further investigations on the association of the neural pathway with respect to the pathophysiology would be helpful to improve our understanding about CP.

## CONFLICTS OF INTEREST

The authors have nothing to disclose.

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# Expression of Human Herpes Virus 6, 7, Epstein-Barr Virus and Cytomegalovirus in Patients with Diverse Adverse Cutaneous Reactions to Drug

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Dear Editor:

Various drugs can cause diverse cutaneous adverse drug reactions (CADR)<sup>1</sup>. Factors have been implicated in CADR, including the dosage, duration of use, physiological status

and genetic background of the patient<sup>1</sup>. In addition, current or past viral infection has been reported to affect the occurrence of CADR<sup>2</sup>. In particular, many authors have suggested the activation of human herpes virus (HHV) 6,

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