Linear and Whorled Nevoid Hypermelanosis with Hemiatrophy

A two-years-old boy presented with abnormal body pigmentation and body asymmetry noted since birth. A preceding vesicular eruption was absent. He was developmentally normal. Family history insignificant. Examination revealed linear and whorled hyperpigmented macules in a reticulate pattern along the Blaschko's lines over trunk and limbs with sparing of palms, soles and mucous membranes (Fig. 1), suggestive of Linear and Whorled Nevoid Hypermelanosis (LWNH). There was left-sided hemi-atrophy. Systemic examination was unremarkable. Fundoscopy was normal. Skin punch biopsy revealed increased pigmentation of the basal layer without incontinence of the pigment. Karyotype, echocardiography and MRI brain were normal.

LWNH is a rare disorder and may be associated with developmental delay, autism, seizures, ocular, cardiac and skeletal abnormalities. Hemi-atrophy has been reported only rarely. It needs to be differentiated from other linear pigmentary disorders distributed along the Blaschko lines. Incontinentia pigmenti is an X-linked



Fig. 1 Linear and Whorled Nevoid Hypermelanosis: Linear and whorled hyperpigmented macules in a reticulate pattern along the Blaschko's lines over trunk and limbs.

dominant disorder presenting in females with a vesicular phase preceding the pigmentary phase. Linear epidermal nevus has characteristic hyperkeratotic plaques. Hypomelanosis of Ito and segmental nevus depigmentosus present with hypopigmented lesions.

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Pityriasis Lichenoides

A 2-year-old male child presented with multiple asymptomatic erythematous scaly papules and plaques with peripheral scales all over the face, scalp, trunk and extremities since 3 weeks (*Fig.*1). Prior to this, there was a history of fluid filled lesions of 2 weeks duration which left behind Chicken pox like scars following symptomatic treatment. Skin biopsy showed typical features of Pityriasis lichenoides et varioliformis acuta (PLEVA). He was treated with narrow band UVB with good results.

Clinical spectrum of Pityriasis lichenoides ranges from acute papular lesions that rapidly evolve into papulovesicles, necrosis and varioliform scarring (PLEVA) to small, scaly, benign-appearing papules (pityriasis lichenoides chronica or PLC) both with a generalized distribution. This affects young adults and occasionally children. Though the etiology of this condition is unclear, infectious/drug related hypersensitivity and premycotic lymphoproliferative disorder are the mainstay theories. The diagnosis is by

histopathology. In PLEVA, as there are polymorphic lesions of papules and vesicles appearing in crops which may heal with scars, a differential diagnosis of varicella has to be considered. Hemorrhagic necrosis and a course of waxing and waning are not common in varicella.



Fig. 1 Erythematous scaly lesions and scars.