A 16-year-old girl presented with asymptomatic, white lesions on the hands and lower extremities. The lesions were first noted five years earlier and had slowly increased in size. There was no family history of a similar skin disorder or autoimmune disease.

Physical examination revealed well-demarcated, chalk-white macules and patches on the dorsum of the hands (Figure 1) and extensor aspects of the legs.

**Question**

Based on the patient’s history and physical examination findings, which one of the following is the most likely diagnosis?

- A. Hypomelanosisis of Ito.
- B. Idiopathic guttate hypomelanosisis.
- C. Pityriasis alba.
- D. Tinea versicolor.
- E. Vitiligo.

See the following page for discussion.
PHOTO QUIZ

Discussion

The answer is E: vitiligo. Vitiligo is characterized by asymptomatic, acquired, amelanotic macules and patches that are milky or chalk-white in color. Lesions are well demarcated, often symmetrical, and show homogenous depigmentation. The most common location is the face, followed by the neck, limbs, and trunk. The onset of vitiligo occurs before 10 years of age in 25% of patients, before 20 years in 50% of patients, and before 40 years in 95% of patients.

In most patients, the disease is slowly progressive, with the appearance of new lesions or enlargement of existing lesions. Vitiligo is more common in those who have a family member with the condition. The cause of vitiligo is unclear, but the patches of depigmented skin are due to the loss of melanocytes.

Hypomelanosis of Ito is characterized by patterned, hypopigmented macules in well-demarcated streaks, stripes, whorls, and patches. It usually involves more than two body segments, most commonly the trunk and proximal extremities. The lesions often present at birth or in the first two years of life and tend to fade in adulthood. Chromosomal abnormalities are found in approximately 50% of cases. About 30% of affected patients have extracutaneous anomalies such as short stature, digital deformity, kyphoscoliosis, psychomotor retardation, hemimegalencephaly, hypotonia, autism, epilepsy, scleral melanocytosis, nystagmus, and strabismus.

Idiopathic guttate hypomelanosis is characterized by multiple asymptomatic, discrete, well-circumscribed, smooth, porcelain-white macules with round or angular borders. Sites of predilection typically include areas with repeated sun exposure, such as the forearms and shins. The condition is more common after 30 years of age. The lesions tend to persist and may slowly enlarge.

Pityriasis alba is characterized by asymptomatic, hypopigmented, round or oval macules. It most commonly affects areas of skin that are rich in sebum production, such as the trunk, upper arms, and neck. Fair-skinned patients tend to have red to brown lesions. Patients with darker skin tend to have hypopigmented lesions. The lesions are covered with a fine scale. The condition is most common among adolescents and young adults. Without antifungal treatment, the disease tends to persist and may slowly enlarge.

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References