Familial generalized lentiginosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial generalized lentiginosis. ORPHA:231040

Familial generalized lentiginosis is a rare, inherited, skin hyperpigmentation disorder characterized by widespread lentigines without associated noncutaneous abnormalities. Patients present multiple brown to dark brown, non-elevated macula of 0.2 to 1 cm in diameter, spread over the entire body, sometimes including palms or soles, but never oral mucosa.