Hashimoto-Pritzker syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hashimoto-Pritzker syndrome. ORPHA:99872

Hashimoto-Pritzker histiocytosis (HPH) is a variant of Langerhans cell histiocytosis (see this term) characterized by multiple disseminated skin lesions (firm, red-brown, painless papulo-nodules).