Schöpf-Schulz-Passarge syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Schöpf-Schulz-Passarge syndrome. ORPHA:50944

Schöpf-Schulz-Passarge syndrome (SSPS) is a rare autosomal recessive ectodermal dysplasia characterized by multiple eyelid apocrine hidrocystomas, palmoplantar keratoderma, hypotrichosis, hypodontia and nail dystrophy.