Diffuse palmoplantar keratoderma-acrocyanosis syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Diffuse palmoplantar keratoderma-acrocyanosis syndrome. ORPHA:86918

Diffuse palmoplantar keratoderma-acrocyanosis syndrome is characterised by the association of diffuse palmoplantar keratoderma and acrocyanosis. It has been described in eight members of one family and in two sporadic cases. The mode of inheritance in the familial cases was autosomal dominant.