Anhidrotic ectodermal dysplasia-immunodeficiency-osteopetrosis-lymphedema syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Anhidrotic ectodermal dysplasia-immunodeficiency-osteopetrosis-lymphedema syndrome. ORPHA:69088

This syndrome is characterized by severe immunodeficiency, osteopetrosis, lymphedema and anhidrotic ectodermal dysplasia.