Sézary syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Sézary syndrome. ORPHA:3162

Sézary syndrome (SS) is an aggressive form of cutaneous T-cell lymphoma characterized by a triad of erythroderma, lymphadenopathy and circulating atypical lymphocytes (Sézary cells).