CHILD syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. CHILD syndrome. ORPHA:139

CHILD syndrome (Congenital Hemidysplasia with Ichthyosiform nevus and Limb Defects, CS) is an X-linked dominant genodermatosis characterized by unilateral inflammatory and scaling skin lesions with ipsilateral visceral and limb anomalies.