Hypertrichosis lanuginosa congenita

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hypertrichosis lanuginosa congenita. ORPHA:2222

Hypertrichosis lanuginosa congenita is a rare congenital skin disease characterized by the presence of 3 to 5cm long lanugo-type hair on the entire body, with the exception of palms, soles, and mucous membranes.