Hypertrichosis cubiti

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hypertrichosis cubiti. ORPHA:2220

Hypertrichosis cubiti is a rare hair anomaly characterized by symmetrical, congenital or early-onset, bilateral hypertrychosis localized on the externsor surfaces of the upper extremities (especially the elbows). Short stature, or other abnormalities, such as developmental delay, facial anomalies and intellectual disability, may or may not be associated.