Hypotrichosis simplex

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hypotrichosis simplex. ORPHA:55654

Hypotrichosis simplex (HS) or hereditary hypotrichosis simplex (HHS) is characterized by reduced pilosity over the scalp and body (with sparse, thin, and short hair) in the absence of other anomalies.