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## Marie Unna hereditary hypotrichosis

**INSERM** 

## Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Marie</u>
<u>Unna hereditary hypotrichosis</u>. ORPHA:444

Marie Unna hereditary hypotrichosis (MUHH) is a rare autosomal dominant hair loss disorder characterized by the absence or scarcity of scalp hair, eyebrows, and eyelashes at birth; coarse and wiry hair during childhood; and progressive hair loss beginning around puberty.

Qeios ID: LA5C6G · https://doi.org/10.32388/LA5C6G