Gingival fibromatosis-hypertrichosis syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Gingival fibromatosis-hypertrichosis syndrome. ORPHA:2026

Gingival fibromatosis - hypertrichosis syndrome is a rare autosomal dominant disorder characterized by a generalized enlargement of the gingiva occurring at birth or during childhood that is associated with generalized hypertrichosis developing at birth, during the first years of life, or at puberty and predominantly affecting the face, upper limbs, and midback.