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# Myhre syndrome

INSERM

## Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base. Myhre syndrome. ORPHA:2588*

Myhre syndrome is characterised by striking muscular build, short stature, reduced joint mobility, brachydactyly, mixed hearing loss and mental retardation of variable severity. Facial dysmorphism with short palpebral fissures, short philtrum, thin lips, maxillary hypoplasia and prognathism is present. Thick skin has been observed in six patients.