Pseudoprogeria syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Pseudoprogeria syndrome. ORPHA:2985

Pseudoprogeria is characterised by intellectual deficit associated with progressive spastic quadriplegia, microcephaly, glaucoma, absence of the eyebrows and eyelashes, and a malformation of the nose. It has been described in two brothers.