Spondyloepiphyseal dysplasia, Nishimura type

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Spondyloepiphyseal dysplasia, Nishimura type. ORPHA:163649

Spondyloepiphyseal dysplasia Nishimura type is characterized by spondyloepiphyseal dysplasia, craniosynostosis, cataracts, cleft palate and intellectual deficit.