Neu-Laxova syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Neu-Laxova syndrome. ORPHA:2671

Neu-Laxova syndrome (NLS) is a rare, multiple malformation syndrome characterised by severe intrauterine growth retardation (IUGR), severe microcephaly with a sloping forehead, severe ichthyosis (collodion baby type), and facial dysmorphism.