# Syndactyly-nystagmus syndrome due to 2q31.1 microduplication

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Syndactyly-nystagmus syndrome due to 2q31.1 microduplication. ORPHA:294026*

A rare, genetic, chromosomal anomaly syndrome resulting from partial duplication of the long arm of chromosome 2 characterized by congenital pendular nystagmus associated with bilateral cutaneous syndactyly between the third and fourth fingers.