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Early-onset non-syndromic cataract

INSERM

Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base*. [Early-onset non-syndromic cataract](#). ORPHA:91492

Early-onset non-syndromic cataract is a rare, genetic, non-syndromic developmental defect of the eye disorder, with high clinical and genetic heterogeneity, most frequently characterized by bilateral, symmetrical, non-progressive cataracts which present at birth or in early-childhood. Additional ocular manifestations (e.g. anterior segment dysgenesis, colobomas, nystagmus, microcornea, microphthalmia, myopia) may be associated, however other organs/systems are usually not affected.