Microcornea-posterior megalolenticonus-persistent fetal vasculature-coloboma syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Microcornea-posterior megalolenticonus-persistent fetal vasculature-coloboma syndrome. ORPHA:231736

Microcornea-posterior megalolenticonus-persistent fetal vasculature-coloboma syndrome is a rare developmental defect of the eye characterized by bilateral microcornea, posterior megalolenticonus, persistent fetal vasculature (extending from the posterior pole of the lens to the optic disc) and posterior chorioretinal coloboma.