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Perlman Syndrome

National Cancer Institute

Source

National Cancer Institute. *Perlman Syndrome*. NCI Thesaurus. Code C103144.

A rare, autosomal recessive syndrome characterized by the presence of polyhydramnios, neonatal macrosomia, craniofacial abnormalities, nephroblastomatosis, and predisposition to Wilms tumor. The prognosis is poor.