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Peters-Plus Syndrome

National Cancer Institute

Source

National Cancer Institute. *Peters-Plus Syndrome*. NCI Thesaurus. Code C123436.

A rare, autosomal recessive inherited syndrome caused by mutations in the B3GALT L gene. It is characterized by abnormalities in the anterior chamber of the eye, short stature, cleft lip with or without cleft palate, distinctive facial features, and intellectual disability.