Peroxisome Biogenesis Disorder 6B

National Cancer Institute

Source

National Cancer Institute. Peroxisome Biogenesis Disorder 6B. NCI Thesaurus. Code C155759.

An autosomal recessive condition caused by mutation(s) in the PEX10 gene, encoding peroxisome biogenesis factor 10. Peroxisome biogenesis disorder 6B is characterized by overlapping phenotypes of neonatal adrenoleukodystrophy and infantile Refsum disease.