Peroxisome Biogenesis Disorder 7B

National Cancer Institute

Source

National Cancer Institute. Peroxisome Biogenesis Disorder 7B. NCI Thesaurus. Code C155761.

An autosomal recessive condition caused by mutation(s) in the PEX26 gene, encoding peroxisome assembly protein 26. Peroxisome biogenesis disorder 7B is characterized by overlapping phenotypes of neonatal adrenoleukodystrophy and infantile Refsum disease.