Peroxisome Biogenesis Disorder 1B

National Cancer Institute

Source

National Cancer Institute. Peroxisome Biogenesis Disorder 1B. NCI Thesaurus. Code C155749.

An autosomal recessive condition caused by mutation(s) in the PEX1 gene, encoding peroxisome biogenesis factor 1. Peroxisome biogenesis disorder 1B is characterized by overlapping phenotypes of neonatal adrenoleukodystrophy and infantile Refsum disease.