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Peroxisome Biogenesis Disorder 5B

National Cancer Institute

Source

National Cancer Institute. <u>Peroxisome Biogenesis Disorder 5B</u>. NCI Thesaurus. Code C155757.

An autosomal recessive condition caused by mutation(s) in the PEX2 gene, encoding peroxisome biogenesis factor 2. Peroxisome biogenesis disorder 5B is characterized by overlapping phenotypes of neonatal adrenoleukodystrophy and infantile Refsum disease.

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