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Peroxisome Biogenesis Disorder 2A

National Cancer Institute

Source

National Cancer Institute. *Peroxisome Biogenesis Disorder 2A*. NCI Thesaurus. Code C155750.

An autosomal recessive condition caused by mutation(s) in the PEX5 gene, encoding peroxisomal targeting signal 1 receptor. Peroxisome biogenesis disorder 2A manifests phenotypically as Zellweger syndrome.