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Familial Paraganglioma-Pheochromocytoma Syndrome

National Cancer Institute

Source

National Cancer Institute. Familial Paraganglioma-Pheochromocytoma Syndrome. NCI Thesaurus. Code C48300.

An autosomal dominant hereditary syndrome caused by mutations in the SDHD, SDHC, SDHB, and SDHAF2 genes. It is characterized by the development of paragangliomas and pheochromocytomas.