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Leber Hereditary Optic Atrophy

National Cancer Institute

Source

National Cancer Institute. *Leber Hereditary Optic Atrophy*. NCI Thesaurus. Code C84808.

A hereditary disorder caused by mitochondrial mutations, resulting in the degeneration of the retinal ganglion cells and optic atrophy. It is characterized by an acute or subacute loss of central vision. It may initially affect one eye only, but eventually the central loss of vision becomes bilateral.