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Waardenburg Syndrome Type 1

National Cancer Institute

Source

National Cancer Institute. *Waardenburg Syndrome Type 1*. NCI Thesaurus. Code C75008.

A rare autosomal dominant syndrome caused by mutations in the PAX3 gene. It is characterized by hearing loss, dystopia canthorum (widely spaced inner corners of the eyes), and changes in the color of the skin, hair, and eyes.