Cone-Rod Dystrophy 2

National Cancer Institute

Source

National Cancer Institute. Cone-Rod Dystrophy 2. NCI Thesaurus. Code C162399.

An inherited condition caused by mutations in the CRX gene, encoding cone-rod homeobox protein. It is characterized by loss of visual acuity in early childhood or late adolescence, impaired color vision, loss of peripheral vision, and nyctalopia. The severity of symptoms may vary.