Revesz Syndrome

National Cancer Institute

Source

National Cancer Institute. Revesz Syndrome. NCI Thesaurus. Code C152064.

An autosomal dominant form of dyskeratosis congenita, caused by mutation(s) in the TINF2 gene, encoding TERF1-interacting nuclear factor 2. It is a fatal disease associated with exudative retinopathy and bone marrow failure.