Mental Retardation, Autosomal Dominant 39

National Cancer Institute

Source

National Cancer Institute. Mental Retardation, Autosomal Dominant 39. NCI Thesaurus. Code C156309.

An autosomal dominant condition caused by mutation(s) in the MYT1L gene, encoding myelin transcription factor 1-like protein. It is characterized by intellectual disability and mild dysmorphic facial features.