MIRAGE Syndrome

National Cancer Institute

Source

National Cancer Institute. MIRAGE Syndrome. NCI Thesaurus. Code C147530.

An autosomal dominant condition caused by mutation(s) in the SAMD9 gene, encoding sterile alpha motif domain-containing protein 9A. It is a syndromic condition comprising myelodysplasia, infection, restriction of growth, adrenal hypoplasia, genital abnormalities, and enteropathy.