Usher Syndrome

National Cancer Institute

Source

National Cancer Institute. Usher Syndrome. NCI Thesaurus. Code C85217.

A rare, autosomal recessive inherited syndrome caused by mutations in the CDH23, CLRN1, GPR98, MYO7A, PCDH15, USH1C, USH1G, and USH2A genes. It is characterized by hearing loss or deafness and progressive loss of vision. The loss of vision is the result of retinitis pigmentosa.