Weill-Marchesani Syndrome

National Cancer Institute

Source

National Cancer Institute. Weill-Marchesani Syndrome. NCI Thesaurus. Code C85226.

A rare, autosomal recessive or dominant inherited connective tissue disorder. The autosomal recessive variant is caused by mutations in the ADAMTS10 gene. It is characterized by abnormalities in the lens of the eye, short stature, brachydactyly, and joint stiffness.