Congenital Contractural Arachnodactyly

National Cancer Institute

Source

National Cancer Institute. Congenital Contractural Arachnodactyly. NCI Thesaurus. Code C129865.

An autosomal dominant connective tissue disorder caused by mutation(s) in the FBN2 gene, encoding fibrillin-2. It is characterized by contractures, arachnodactyly, scoliosis, micrognathia, and crumpled ears.