Spinal Muscular Atrophy Type 2

National Cancer Institute

Source

National Cancer Institute. Spinal Muscular Atrophy Type 2. NCI Thesaurus. Code C156310.

An autosomal recessive condition caused by mutation(s) in the SMN1 gene, encoding survival motor neuron protein. It is characterized by onset between 3 and 15 months of age, and is intermediate in terms of severity between spinal muscular atrophy (SMA) type I and SMA type III.