Familial Hypertrophic Cardiomyopathy Type 2

National Cancer Institute

Source

National Cancer Institute. Familial Hypertrophic Cardiomyopathy Type 2. NCI Thesaurus. Code C142892.

An autosomal dominant subtype of familial hypertrophic cardiomyopathy caused by mutation(s) in the TNNT2 gene, encoding troponin T, cardiac muscle.