Carboxypeptidase N Deficiency

National Cancer Institute

Source

National Cancer Institute. Carboxypeptidase N Deficiency. NCI Thesaurus. Code C132196.

An autosomal recessive condition caused by mutation(s) in the CPN1 gene, encoding carboxypeptidase N catalytic chain. It may be characterized by episodic angioedema, chronic urticaria, asthma and/or allergic hypersensitivity.