SNX29 wt Allele

National Cancer Institute

Source

National Cancer Institute. SNX29 wt Allele. NCI Thesaurus. Code C97765.

Human SNX29 wild-type allele is located in the vicinity of 16p13.13 and is approximately 598 kb in length. This allele, which encodes sorting nexin-29 protein, may play a role in the regulation of GTPase-mediated signaling or lipid trafficking. A point mutation in the gene may be associated with ovarian serous carcinoma.