Resource Summary Report

Generated by RRID on Aug 24, 2026

pCDNA3-Flag-SPRTN-Y117C

RRID:Addgene_110216
Type: Plasmid

Proper Citation

RRID:Addgene_110216

Plasmid Information

URL: http://www.addgene.org/110216

Proper Citation: RRID:Addgene_110216

Insert Name: SPRTN

Organism: Homo sapiens

Bacterial Resistance: Ampicillin

Defining Citation: PMID:27871366

Vector Backbone Description: Backbone Marker:Invitrogen; Vector Backbone:pCDNA3.1; Vector Types:Mammalian Expression; Bacterial Resistance:Ampicillin

Comments: According to the large datasets deposited in various databses like: dbSNP, 1000 Genomes, the ExAC and the gnomAD browser, the variant c.887C>T, p.Pro296Leu is found roughly in 50% of general world population. Moreover, according to the large-scale sequencing studies involving more than 135 000 healthy individuals deposited in The Genome Aggregation Database (gnomAD), it is more prevalent than the previously assigned reference allele in the European population, with a minor allele frequency of 0.69: http://gnomad.broadinstitute.org/variant/1-231488524-C-T Taken this into account, this variant (P296L) is s actually the accurate reference (wild-type) allele, and thus this variant can not be the cause of any Mendelian disease. This is the reason why all our vectors bear this variant.

Plasmid Name: pCDNA3-Flag-SPRTN-Y117C

Relevant Mutation: Y117C, P296L (see depositor comments below)

Record Creation Time: 20220422T221536+0000

Record Last Update: 20260815T080138+0000

Ratings and Alerts

No rating or validation information has been found for pCDNA3-Flag-SPRTN-Y117C.

No alerts have been found for pCDNA3-Flag-SPRTN-Y117C.

Data and Source Information

Source: Addgene

Usage and Citation Metrics

We have not found any literature mentions for this resource.