

OriGene Technologies, Inc.

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Product datasheet for TP320271

PPP2R2B (NM_004576) Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Description: Purified recombinant protein of Homo sapiens protein phosphatase 2 (formerly 2A),

regulatory subunit B, beta isoform (PPP2R2B), transcript variant 1

Species: Human Expression Host: HEK293

Tag: C-Myc/DDK

Predicted MW: 51.5 kDa

Concentration: >50 ug/mL as determined by microplate BCA method

Purity: > 80% as determined by SDS-PAGE and Coomassie blue staining

Buffer: 25 mM Tris.HCl, pH 7.3, 100 mM glycine, 10% glycerol.

Preparation: Recombinant protein was captured through anti-DDK affinity column followed by

conventional chromatography steps.

Storage: Store at -80°C.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and

handling conditions. Avoid repeated freeze-thaw cycles.

RefSeq: NP 004567

Locus ID:5521RefSeq Size:2300Cytogenetics:5q32RefSeq ORF:1329

Synonyms: B55BETA; PP2AB55BETA; PP2ABBETA; PP2APR55B; PP2APR55BETA; PR2AB55BETA;

PR2ABBETA: PR2APR55BETA: PR52B





Summary:

The product of this gene belongs to the phosphatase 2 regulatory subunit B family. Protein phosphatase 2 is one of the four major Ser/Thr phosphatases, and it is implicated in the negative control of cell growth and division. It consists of a common heteromeric core enzyme, which is composed of a catalytic subunit and a constant regulatory subunit, that associates with a variety of regulatory subunits. The B regulatory subunit might modulate substrate selectivity and catalytic activity. This gene encodes a beta isoform of the regulatory subunit B55 subfamily. Defects in this gene cause autosomal dominant spinocerebellar ataxia 12 (SCA12), a disease caused by degeneration of the cerebellum, sometimes involving the brainstem and spinal cord, and in resulting in poor coordination of speech and body movements. Multiple alternatively spliced variants, which encode different isoforms, have been identified for this gene. The 5' UTR of some of these variants includes a CAG trinucleotide repeat sequence (7-28 copies) that can be expanded to 55-78 copies in cases of SCA12. [provided by RefSeq, Jul 2016]

Protein Families: Druggable Genome, Phosphatase

Protein Pathways: Tight junction