

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

linkedin.com/company/szaboscandic in





PPP2R2B (Human) IP-WB Antibody Pair

Catalog #: H00005521-PW1 規格:[1 Set]

List All

Specification		Application Image
Product Description:	This IP-WB antibody pair set comes with one antibody for immunoprecipitation and another to detect the precipitated protein in western blot.	Immunoprecipitation-Westerr Blot
Reactivity:	Human	
Quality Control Testing:	Immunoprecipitation-Western Blot (IP-WB) 250 =	
	Immunoprecipitation of PPP2R2B transfected lysate using mouse monoclonal anti-PPP2R2B and Protein A Magnetic Bead (<u>U0007</u>), and immunoblotted with rabbit polyclonal anti-PPP2R2B.	
Supplied Product:	Antibody pair set content: 1. Antibody pair for IP: mouse monoclonal anti-PPP2R2B (300 ug) 2. Antibody pair for WB: rabbit polyclonal anti-PPP2R2B (50 ul)	
Storage Instruction:	Store reagents of the antibody pair set at -20°C or lower. Please aliquot to avoid repeated freeze thaw cycle. Reagents should be returned to -20°C storage immediately after use.	
MSDS:	<u>Download</u>	
Applications		
Immunoprecipit	ation-Western Blot	
Gene Information	on	
Entrez GeneID:	<u>5521</u>	
Gene Name:	PPP2R2B	
Gene Alias:	B55-BETA,FLJ95686,MGC24888,PP2A-B55BETA,PP2A-PR55B,PP2AB-BETA,PP2APR55-BETA,PR2AB-BETA,PR2AB55-BETA,PR2APR55-BETA,PR52B,PR55-BETA,SCA12	
Gene Description:	protein phosphatase 2 (formerly 2A), regulatory subunit B, beta isoform	
Omim ID:	<u>604325, 604326</u>	
	D 4 60	

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Gene Ontology: Hyperlink

Gene 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 66-78 copies in cases of SCA12. [provided by RefSeq

Other Designations:

PP2A, subunit B, B-beta isoform, PP2A, subunit B, R2-beta isoform, beta isoform of regulatory subunit B55, protein phosphatase 2, protein phosphatase 2 (formerly 2A), regulatory subunit B (PR 52), beta isoform, serine/threonine protein phosphatase 2A, 55 kDa

Gene Pathway

Tight junction

Related Disease

Alzheimer Disease Alzheimer disease Cardiovascular Diseases Cerebellar Ataxia <u>Chronic Disease Diabetes Mellitus, Type 2 Disease Progression Edema Essential Tremor</u> Essential tremor Genetic Predisposition to Disease Genomic Instability Parkinson disease Schizophrenia Schizophrenia Spinocerebellar ataxia Spinocerebellar Ataxias Tobacco Use Disorder

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