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anti-FKBP1A antibody (N-Term)

2 Images



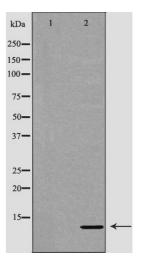
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Overview

100 μL
FKBP1A
N-Term
Human, Mouse, Rat
Rabbit
Polyclonal
Western Blotting (WB), Immunohistochemistry (IHC), ELISA
A synthesized peptide derived from human FKBP1A, corresponding to a region within N-
terminal amino acids.
IgG
FKBP1A Antibody detects endogenous levels of total FKBP1A.
Pig,Bovine,Rabbit,Xenopus
The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling
Resin (Thermo Fisher Scientific).
FKBP1A

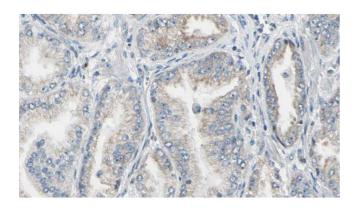
Target Details

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Background:	Description: Keeps in an inactive conformation TGFBR1, the TGF-beta type I serine/threonine kinase receptor, preventing TGF-beta receptor activation in absence of ligand. Recruits SMAD7 to ACVR1B which prevents the association of SMAD2 and SMAD3 with the activin receptor complex, thereby blocking the activin signal. May modulate the RYR1 calcium channel activity PPIases accelerate the folding of proteins. It catalyzes the cis-trans isomerization of proline imidic peptide bonds in oligopeptides. Gene: FKBP1A
Molecular Weight:	12kDa
Gene ID:	2280
UniProt:	P62942
Pathways:	Negative Regulation of Transporter Activity, Methionine Biosynthetic Process
Application Details	
Application Notes:	WB 1:500-1:2000, IHC 1:50-1:200, ELISA(peptide) 1:20000-1:40000
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Concentration:	1 mg/mL
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C
Storage Comment:	Store at -20 °C. Stable for 12 months from date of receipt.
Expiry Date:	12 months



Western Blotting

Image 1. Western blot analysis of extracts of mouse cerebellum tissue, using FKBP1A antibody. The lane on the left is treated with the antigen-specific peptide.



Immunohistochemistry

Image 2. ABIN6276862 at 1/100 staining Human prostate tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the antibody for 1.5 hours at 22¡ãC. An HRP conjugated goat anti-rabbit antibody was used as the secondary