Revision 5	
AMPA Receptor 4 (GluA 4) (D41A11) XP <sup>®</sup> Rabbit mAb	<b>Cell Signaling</b> TECHNOLOGY®
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Applications: N, W-S, IP, IF-F, IF- IC	<b>Reactivity:</b> H M R	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 100	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #P48058	Entrez-Gene Io 2893	
Product Usage Information		Application Western Blotting Simple Western™ Immunoprecipitation Immunofluorescence				<b>Dilution</b> 1:1000 1:10 - 1:50 1:50 1:400	
Storage		Immunofluorescence (Immunocytochemistry) 1:400 Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less th					
Storage	0.02% sodium azide. Store at -20°C. Do not aliquot the antibody.						
		For a carrier free (BSA and azide free) version of this product see product #26485.					
Specificity/Sensi	tivity	AMPA Receptor 4 (GluA 4) (D41A11) XP $^{ m (B}$ Rabbit mAb detects endogenous levels of total GluA 4 protein.					
Source / Purifica	tion	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Gln890 of human GluA 4 protein.					
Background		AMPA- (α-amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid), kainate-, and NMDA- (N-methyl-D- aspartate) receptors are the three main families of ionotropic glutamate-gated ion channels. AMPA receptors (AMPARs) are comprised of four subunits (GluR 1-4), which assemble as homo- or hetero- tetramers to mediate the majority of fast excitatory transmissions in the central nervous system. AMPARs are implicated in synapse formation, stabilization, and plasticity (1). In contrast to GluR 2- containing AMPARs, AMPARs that lack GluR 2 are permeable to calcium (2). Post-transcriptional modifications (alternative splicing, nuclear RNA editing) and post-translational modifications (glycosylation, phosphorylation) result in a very large number of permutations, fine-tuning the kinetic properties of AMPARs. Research studies have implicated activity changes in AMPARs in a variety of diseases including Alzheimer's, amyotrophic lateral sclerosis (ALS), stroke, and epilepsy (1).					
				ynapses and GluR 4 deli orylation of Ser842 by PK		nd cell surface	
Background Ref	erences	1. Palmer, C.L. et al. (2005) <i>Pharmacol Rev</i> 57, 253-77. 2. Cull-Candy, S. et al. (2006) <i>Curr Opin Neurobiol</i> 16, 288-97. 3. Gomes, A.R. et al. (2007) <i>Traffic</i> 8, 259-69.					
Species Reactivi	ty	Species reactivity is de	etermined by testin	g in at least one approve	ed application (e.g.	., western blot).	
Western Blot Bu	ffer	IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.					
Applications Key	1	W: Western Blotting W-S: Simple Western™ IP: Immunoprecipitation IF-F: Immunofluorescence (Frozen) IF-IC: Immunofluorescence (Immunocytochemistry)					
Cross-Reactivity	Кеу	H: Human M: Mouse R: Rat					
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