[Primary Antibody]

DRD1 Recombinant Rabbit mAb



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- DATASHEFT		400-901-9800	
Host: Rabbit	Isotype: IgG	Applications: WB (1:500-1000)	
Clonality: Recombinant	CloneNo.: 3C3	IHC-P (1:50-200) IHC-F (1:50-200) IF (1:50-200)	
GenelD: 1812	SWISS: P21728		
Target: DRD1		ICC/IF (1:50)	
Purification: affinity purified by Protein A		Reactivity: Human, Mouse, Rat	
Concentration: 1mg/ml			
Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.		Predicted MW.: ^{50 kDa}	
Background: This gene encodes the D1 subtype of the dopamine receptor. The D1 subtype is the most abundant dopamine receptor in the central nervous system. This G-protein coupled receptor stimulates adenylyl cyclase and activates cyclic AMP-dependent protein kinases. D1 receptors regulate neuronal growth and development, mediate some behavioral responses, and modulate dopamine receptor D2-mediated events. Alternate transcription initiation sites result in two transcript variants of this gene. [provided by RefSeq, Jul 2008]		Subcellular Location: Cell membrane ,Cytoplasm	

– VALIDATION IMAGES



Sample: Lane 1: Human SH-SY5Y cell lysates Primary: Anti-DRD1 (bsm-52920R) at 1/200 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 50 kDa Observed band size: 70 kDa



Sample: Lane 1: mouse kidney cells lysates Primary: Anti-Dopamine Receptor D1 (bsm-52920R) at 1:500 dilution Secondary: Goat Anti-Rabbit IgG - HRP at 1:5000 dilution Predicted band size: 50 kD Observed band size: 75 kD



Paraformaldehyde-fixed, paraffin embedded (Human kidney); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Dopamine Receptor D1) Monoclonal Antibody, Unconjugated (bsm-52920R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Paraformaldehyde-fixed, paraffin embedded (mouse brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Dopamine Receptor D1) Monoclonal Antibody, Unconjugated



SH-SY5Y cell; 4% Paraformaldehyde-fixed; Triton X-100 at room temperature for 20 min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Antibody incubation with (DRD1) monoclonal Antibody, Unconjugated (bsm-52920R) 1:50, 90 minutes at 37°C; followed by a conjugated Goat Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002)



N2A cell; 4% Paraformaldehyde-fixed; Triton X-100 at room temperature for 20 min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Antibody incubation with (DRD1) monoclonal Antibody, Unconjugated (bsm-52920R) 1:50, 90 minutes at 37°C; followed by a conjugated Goat Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002)

- SELECTED CITATIONS ------

• [IF=17.694] Carpenter, Marco D.. et al. Cell-type specific profiling of histone post-translational modifications in the adult mouse striatum. NAT COMMUN. 2022 Dec;13(1):1-12 IHC ;Mouse. 36513652