

3626

DAG Lipase α (D3G8H) Rabbit mAb



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Applications: W, IP	Reactivity: H M R	Sensitivity: Endogenous	MW (kDa): 115	Source/Isotype: Rabbit IgG	UniProt ID: #Q9Y4D2	Entrez-Gene Id: 747
Product Usage Information		Application Western Blotting Immunoprecipitation			Dilution 1:1000 1:50	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		DAG Lipase α (D3G8H) Rabbit mAb recognizes endogenous levels of total DAG lipase α protein. This antibody is not predicted to cross-react with DAG lipase β based on sequence homology of the antigen.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ser791 of human DAG lipase α protein.				
Background		Diacylglycerol (DAG) lipases comprise two enzymes called DAG lipase α and β , which are the products of two related genes (1). DAG lipases are transmembrane proteins composed of a short amino-terminal intracellular domain, four transmembrane domains, and a large carboxy-terminal cytoplasmic domain containing the active site. These enzymes are responsible for the biosynthesis of 2-acylglycerol from diacylglycerol in a calcium-dependent manner (1). One of the major endocannabinoid ligands that activate cannabinoid receptors, 2-arachidonyl glycerol (2-AG), is produced by DAG lipases (2). Research studies suggest that DAG lipase α is the isoform primarily responsible for the central production of 2-AG (3). DAG lipase β has been implicated in studies of 2-AG production at the periphery in specific cell types and pathophysiological contexts, such as in hepatic stellate cells during alcohol induced fatty liver (4).				
Background References		1. Bisogno, T. et al. (2003) <i>J Cell Biol</i> 163, 463-8. 2. Mechoulam, R. et al. (1995) <i>Biochem Pharmacol</i> 50, 83-90. 3. Yoshino, H. et al. (2011) <i>J Physiol</i> 589, 4857-84. 4. Jeong, W.I. et al. (2008) <i>Cell Metab</i> 7, 227-35.				
Species Reactivity		Species reactivity is determined by testing in at least one approved application (e.g., western blot).				
Western Blot Buffer		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		W: Western Blotting IP: Immunoprecipitation				
Cross-Reactivity Key		H: Human M: Mouse R: Rat				
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