GM130 (D6B1) XP® Rabbit mAb



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Applications: W, W-S, IP, IF-IC	Reactivity: H Mk	Sensitivity: Endogenous	MW (kDa): 140	Source/Isotype: Rabbit IgG	UniProt ID: #Q08379	Entrez-Gene Id: 2801
Product Usage Information		Application Western Blotting Simple Western™ Immunoprecipitation Immunofluorescence (Immunocytochemistry)			Dilution 1:1000 1:50 - 1:250 1:100 1:3200	
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		GM130 (D6B1) XP [®] Rabbit mAb recognizes endogenous levels of total GM130 protein. This antibody may cross-react with a protein of unknown origin at 30 kDa.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Thr185 of human GM130 protein.				
Background		The Golgi apparatus functions in the modification, organization, and transport of proteins and membranes targeted to other parts of the cell, such as the plasma membrane, lysosomes, and endosomes. This regulated transport is important for appropriate protein localization, secretion, and signal transduction. Members of the Golgin family of proteins, including GM130, Giantin, p115, and GRASP65, are defined by their presence in the Golgi matrix and by their long coiled-coil domains. Golgin function, which is regulated in part by small GTPases of the Rab and Arl families, includes establishing and maintaining Golgi structure and transport (reviewed in 1). The Golgi cisternae are stacked and linked laterally to form a ribbon. GRASP65 and GM130 are required for membrane fusion events that mediate ribbon formation during Golgi assembly. These lateral fusion events allow for uniform distribution of Golgi enzymes (2). GM130 and Giantin interact with the transport factor p115 to facilitate endoplasmic reticulum (ER)-Golgi transport (3). GM130 is also involved in the transport of the ether-a-go-go-related (hERG) potassium ion channel. Inappropriate hERG localization may be an underlying cause of long QT syndrome, a hereditary and potentially fatal cardiac arrhythmia (4). Further, GM130 was implicated in signal transduction regulating invasion, migration, and cell polarization via its interaction with and activation of serine/threonine kinases YSK1 and Mst4 (5).				
Background References		1. Barr, F.A. and Short, B. (2003) <i>Curr. Opin. Cell Biol.</i> 15, 405-413. 2. Puthenveedu, M.A. et al. (2006) <i>Nat. Cell Biol.</i> 8, 238-248. 3. Alvarez, C. et al. (2001) <i>J. Biol. Chem.</i> 276, 2693-2700. 4. Roti, E.C. et al. (2002) <i>J. Biol. Chem.</i> 277, 47779-47785. 5. Preisinger, C. et al. (2004) <i>J. Cell Biol.</i> 164, 1009-1020.				
Species Reactiv	ity	Species reactivity is de	etermined by testing	g in at least one approve	ed application (e.g.,	western blot).
Western Blot Buffer		IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.				

Applications Key

W: Western Blotting **W-S**: Simple Western™ **IP**: Immunoprecipitation **IF-IC**: Immunofluorescence (Immunocytochemistry)

Cross-Reactivity Key

H: Human Mk: Monkey

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