

Sec31A (D1G7I) Rabbit mAb



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Applications: W, IF-IC	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 95-140	Source/Isotype: Rabbit IgG	UniProt ID: #O94979	Entrez-Gene Id: 22872
Product Usage Information		Application Western Blotting Immunofluorescence			# 69.137 3	Dilution 1:1000 1:400
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		Sec31A (D1G7I) Rabbit mAb recognizes endogenous levels of total Sec31A protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ser86 of human Sec31A protein.				
Background		The coat protein complex II (COPII) is composed of five cytosolic proteins and includes the Sec23/24 complex, the Sec13/31 complex, and Sar1. The COPII coat is located at the ER/Golgi interface and is involved in transport of newly synthesized proteins from the ER to the Golgi apparatus (1). COPII formation is initiated through the binding of the activated G protein, Sar1, to the Sec23/24 complex to form a pre-budding complex that directly binds target molecules (1-3). This pre-budding complex further recruits Sec13/31 to form mature COPII coat (4,5). The Sec31 subunit of COPII coat interacts with Sec13 at the ER exit and is required for both vesicle formation and ER-Golgi transport. Two isoforms of human Sec31 have been identified, Sec31A and Sec31B, which share a sequence homology of 47.3% (6-8). Sec31A is ubiquitously expressed in tissues and organs, whereas Sec31B is enriched in brain and testis (7,8). In classical Hodgkin lymphoma, a novel fusion of Jak2 with Sec31A renders Jak2 constitutively active and subject to Jak2 inhibitor effects (9).				
Background References		1. Aridor, M. et al. (1998) <i>J Cell Biol</i> 141, 61-70. 2. Miller, E.A. et al. (2003) <i>Cell</i> 114, 497-509. 3. Mossessova, E. et al. (2003) <i>Cell</i> 114, 483-95. 4. Barlowe, C. et al. (1994) <i>Cell</i> 77, 895-907. 5. Bi, X. et al. (2007) <i>Dev Cell</i> 13, 635-45. 6. Shugrue, C.A. et al. (1999) <i>J Cell Sci</i> 112 (Pt 24), 4547-56. 7. Tang, B.L. et al. (2000) <i>J Biol Chem</i> 275, 13597-604. 8. Stankewich, M.C. et al. (2006) <i>J Cell Sci</i> 119, 958-69. 9. Van Roosbroeck, K. et al. (2011) <i>Blood</i> 117, 4056-64.				

Species Reactivity Species react

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 nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key W: Western Blotting IF

 $\textbf{W:} \ \textbf{Western Blotting IF-IC:} \ \textbf{Immunofluorescence (Immunocytochemistry)}$

Cross-Reactivity Key H: Human M: Mouse R: Rat Mk: Monkey

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