## **.5601**

## βIG-H3 (D31B8) XP® Rabbit mAb



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## For Research Use Only. Not for Use in Diagnostic Procedures.

<b>Applications:</b> W, IP, IF-IC	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 70	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #Q15582	Entrez-Gene Id: 7045
Product Usage Information		<b>Application</b> Western Blotting Immunoprecipitation Immunofluorescence	(Immunocytochem	istry)		<b>Dilution</b> 1:1000 1:100 1:800
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		$\beta IG\text{-H3}$ (D31B8) XP $^{\! @}$ Rabbit mAb detects endogenous levels of total $\beta IG\text{-H3}$ protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human $\beta$ IG-H3 protein.				
Background		$\beta$ IG-H3 (TGFBI/RGD-CAP/Kerato-epithelin) is a 683-amino acid secretory protein induced by TGF- $\beta$ that plays a role in cell adhesion, differentiation, and apoptosis (1-4). $\beta$ IG-H3 contains an internal cysteinerich EMI domain followed by four fasciclin-1 domains and a carboxy terminal RGD domain (1,2). It contributes to cell adhesion through interactions with integrins as well as a number of extracellular matrix (ECM) proteins including collagen, fibronectin, and laminin (5-7). ECM $\beta$ IG-H3 is found in a wide variety of tissues (8-12). Mutations in the $\beta$ IG-H3 gene as well as elevated protein levels are most notably associated with corneal dystrophies (13).				
Background Ro	eferences	1. Skonier, J. et al. (1992) <i>DNA Cell Biol</i> 11, 511-22. 2. Skonier, J. et al. (1994) <i>DNA Cell Biol</i> 13, 571-84. 3. Hashimoto, K. et al. (1997) <i>Biochim Biophys Acta</i> 1355, 303-14. 4. Kim, J.E. et al. (2003) <i>Oncogene</i> 22, 2045-53. 5. Kim, J.E. et al. (2002) <i>Invest Ophthalmol Vis Sci</i> 43, 656-61. 6. Billings, P.C. et al. (2002) <i>J Biol Chem</i> 277, 28003-9. 7. Hanssen, E. et al. (2003) <i>J Biol Chem</i> 278, 24334-41. 8. Gibson, M.A. et al. (1997) <i>J Histochem Cytochem</i> 45, 1683-96. 9. Billings, P.C. et al. (2000) <i>Am J Respir Cell Mol Biol</i> 22, 352-9. 10. Gilbert, R.E. et al. (1998) <i>Kidney Int</i> 54, 1052-62. 11. Rawe, I.M. et al. (1997) <i>Invest Ophthalmol Vis Sci</i> 38, 893-900. 12. LeBaron, R.G. et al. (1995) <i>J Invest Dermatol</i> 104, 844-9. 13. Munier, F.L. et al. (1997) <i>Nat Genet</i> 15, 247-51.				

**Species Reactivity** Species reactivity is determined

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 IF-IC: Immunofluorescence (Immunocytochemistry)

Cross-Reactivity Key H: Human

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