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-20C FGF19 (D1N3R) Rabbit mAb #83348



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Applications: Reactivity: W, IHC-P H	Sensitivity: Endogenous	MW (kDa): 22	Source/Isotype: Rabbit IgG	UniProt ID: #O95750	Entrez-Gene Id: 9965	
Product Usage Information	Application Western Blotting Immunohistochemistry (Paraffin)			Dilution 1:1000 1:200		
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 aliguot the antibody.					
Specificity/Sensitivity	FGF19 (D1N3R) Rabbit mAb recognizes endogenous levels of total human FGF19 protein. This antibody does not cross-react with the mouse ortholog FGF15.					
Source / Purification	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ala148 of human FGF19 protein.					
Background	FGF19 is a member of the large and diverse Fibroblast Growth Factor (FGF) family of peptide growth factors. FGF19 functions as a high affinity ligand for the FGF4 receptor, to which it binds in a heparin- dependent manner (1). Under normal physiological conditions, FGF19 is produced in the ileum when absorbed bile acids bind to the farnesoid X receptor (FXR), activating transcription of the FGF19 gene. FGF19 in turn functions in a negative feedback fashion to regulate bile acid synthesis (2). Consequently, disruptions in FGF19 signaling have been linked with clinical defects in bile acid synthesis, which may manifest as primary bile acid diarrhea or cholestasis (3,4). Research studies in oncology have shown that FGF19 can function in an endocrine, paracrine or autocrine fashion to promote tumorigenesis. In human studies, this has been demonstrated for breast cancer (5), gastric cancer (6) prostate cancer (7, 8), and hepatocellular carcinoma (9). Similar tumorigenic effects have been described for the mouse ortholog (Fgf15), notably in genetic models of hepatocellular carcinoma (10, 11)					
Background References	 Xie, M.H. et al. (1999) <i>Cytokine</i> 11, 729-35. Miyata, M. et al. (2012) <i>J Steroid Biochem Mol Biol</i> 132, 41-7. Walters, J.R. et al. (2009) <i>Clin Gastroenterol Hepatol</i> 7, 1189-94. Schaap, F.G. et al. (2009) <i>Hepatology</i> 49, 1228-35. Tiong, K.H. et al. (2016) Oncotarget 7, 57633-50. Wang, S. et al. (2016) <i>Oncol Res</i> 23, 197-203. Nagamatsu, H. et al. (2015) <i>Prostate</i> 75, 1092-101. Feng, S. et al. (2013) <i>Cancer Res</i> 73, 2551-62. Sawey, E.T. et al. (2011) <i>Cancer Cell</i> 19, 347-58. Desnoyers, L.R. et al. (2008) <i>Oncogene</i> 27, 85-97. Uriarte, I. et al. (2015) <i>Int J Cancer</i> 136, 2469-75. 					
Species Reactivity	Species reactivity is d	etermined by testin	g in at least one approve	ed application (e.g.	western blot)	
Western Blot Buffer	Species reactivity is determined by testing in at least one approved application (e.g., western blot). 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 IHC-P: Immunohistochemistry (Paraffin)					
Cross-Reactivity Key	H: Human					
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