– DATASHEET –

[Primary Antibody]

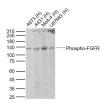
phospho-FGFR1 (Tyr154) Rabbit pAb



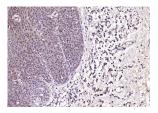
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- DATASHEET		
Host: Rabbit	lsotype: IgG	Applications: WB (1:500-2000)
Clonality: Polyclonal		IHC-P (1:100-500) IHC-F (1:100-500)
GenelD: 2260	SWISS: P11362	IF (1:100-500)
Target: FGFR1 (Tyr154)		Reactivity: Human, Mouse
Immunogen: KLH conjugated Synthesised phosphopeptide derived from human FGFR1 around the phosphorylation site of Tyr154: AP(p-Y)WT.		(predicted: Rat, Rabbit, Pig, Sheep, Cow, Dog, GuineaPig, Horse)
Purification: affinity purified by Protein A		
Concentration: 1mg/ml		Predicted MW.: ^{88 kDa}
Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.		Subcellular Cell membrane ,Cytoplasm Location: ,Nucleus
Background: Fibroblast growth factors (FGFs) produce mitogenic and angiogenic effects in target cells by signaling through the cellular surface tyrosine kinase receptors. There are four members of the FGF receptor family: FGFR-1 (flg), FGFR-2 (bek, KGFR), FGFR-3 and FGFR-4. Each receptor contains an extracellular ligand binding domain, a transmembrane region and a cytoplasmic kinase domain (1). Following ligand binding and dimerization, the receptors are phosphorylated at specific tyrosine residues (2). Seven tyrosine residues in the cytoplasmic tail of FGFR-1 can be phosphorylated: Tyr463, Tyr583, Tyr585, Tyr653, Tyr654, Tyr730 and Tyr766. Tyrosine 653 and 654 are important for catalytic activity of the activated FGFR and are essential for signaling (3). The other phosphorylated tyrosine residues may provide docking sites for downstream signaling components such as Crk and PLCgamma.		

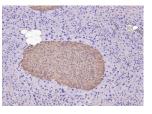
– VALIDATION IMAGES



Sample: Lane 1: Human A673 cell lysates Lane 2: Human A431 cell lysates Lane 3: Human Molt-4 cell lysates Lane 4: Human U87MG cell lysates Primary: Anti-Phospho-FGFR1 (Tyr154) (bs-13155R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 88 kD Observed band size: 130 kD



Paraformaldehyde-fixed, paraffin embedded (human laryngeal carcinoma); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Phospho-FGFR1 (Tyr154)) Polyclonal Antibody, Unconjugated (bs-13155R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Paraformaldehyde-fixed, paraffin embedded (mouse pancreas); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Phospho-FGFR1 (Tyr154)) Polyclonal Antibody, Unconjugated (bs-13155R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.

- SELECTED CITATIONS -

• [IF=6.353] Song Siqi. et al. Specific bFGF targeting of KIM-1 in ischemic kidneys protects against renal ischemia reperfusion injury in rats. REGEN BIOMATER. 2022 May:: WB ;Human,Rat. 35615568