[Primary Antibody]

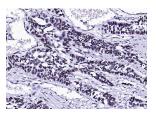
phospho-FGFR1 (Tyr307) Rabbit pAb



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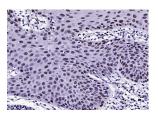
— DATASHEET – Host: Rabbit Isotype: IgG Applications: IHC-P (1:100-500) IHC-F (1:100-500) Clonality: Polyclonal IF (1:100-500) GenelD: 2260 SWISS: P11362 ELISA (1:5000-10000) Target: FGFR1 (Tyr307) Reactivity: Human, Rat, Rabbit Immunogen: KLH conjugated Synthesised phosphopeptide derived from human (predicted: Mouse, Pig. FGFR1 around the phosphorylation site of Tyr307: LP(p-Y)VQ. < Cow, Chicken, GuineaPig, Horse) Extracellular > Purification: affinity purified by Protein A Predicted 90 kDa MW.: Concentration: 1mg/ml Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Subcellular Cell membrane, Cytoplasm Glycerol. Location: ,Nucleus Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles. 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).

- VALIDATION IMAGES



PLCgamma.

Paraformaldehyde-fixed, paraffin embedded (Human esophageal cancer); 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 (Tyr307)) Polyclonal Antibody, Unconjugated (bs-5325R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



The other phosphorylated tyrosine residues may provide docking sites for downstream signaling components such as Crk and

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 (Tyr307)) Polyclonal Antibody, Unconjugated (bs-5325R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.