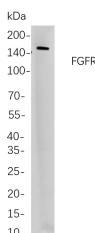


Recombinant FGFR1 Monoclonal Antibody

catalog number: AN301100L

Note: Centrifuge before opening to ensure complete recovery of vial contents.

| Description | |
|--|---|
| Reactivity | Human |
| Immunogen | Recombinant Human FGFR1 protein |
| Host | Rabbit |
| Isotype | IgG,κ |
| Clone | B855 |
| Purification | Protein A |
| Buffer | PBS, 50% glycerol, 0.05% Proclin 300, 0.05% protein protectant. |
| Applications | Recommended Dilution |
| IHC | 1:100-200 |
| WB | 1:500-2000 |
| IF | 1:200-1000 |
| ELISA | 1:5000-20000 |
| IP | 1:50-200 |
| Data | |
|  | |
| <p>Western Blot with Recombinant FGFR1 Monoclonal Antibody at dilution of 1:1000 dilution. Lane A: HEK293 cells.</p> <p>Observed-MW:145 kDa Calculated-MW:91 kDa</p> | |
| Preparation & Storage | |
| Storage | Store at -20°C Valid for 12 months. Avoid freeze / thaw cycles. |
| Shipping | Ice bag |
| Background | |

For Research Use Only

Toll-free: 1-888-852-8623

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Tel: 1-832-243-6086

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Rev. V1.2

The protein encoded by this gene is a member of the fibroblast growth factor receptor (FGFR) family, where amino acid sequence is highly conserved between members and throughout evolution. FGFR family members differ from one another in their ligand affinities and tissue distribution. A full-length representative protein consists of an extracellular region, composed of three immunoglobulin-like domains, a single hydrophobic membrane-spanning segment and a cytoplasmic tyrosine kinase domain. The extracellular portion of the protein interacts with fibroblast growth factors, setting in motion a cascade of downstream signals, ultimately influencing mitogenesis and differentiation. This particular family member binds both acidic and basic fibroblast growth factors and is involved in limb induction. Mutations in this gene have been associated with Pfeiffer syndrome, Jackson-Weiss syndrome, Antley-Bixler syndrome, osteoglophonic dysplasia, and autosomal dominant Kallmann syndrome 2. Chromosomal aberrations involving this gene are associated with stem cell myeloproliferative disorder and stem cell leukemia lymphoma syndrome. Alternatively spliced variants which encode different protein isoforms have been described; however, not all variants have been fully characterized.

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