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Product Information

MemDX™ Membrane Protein Human FGFR2 (Fibroblast growth factor receptor 2) expressed in Sf9 for Antibody Discovery

Cat. No.: **MP0131Q**
This product is for research use only and is not intended for diagnostic use.

This product is a 39.5 kDa Human FGFR2 membrane protein expressed in Sf9. The protein is for research use only and is not approved for use in humans or in clinical diagnosis.

Product Specifications

Host Species
Human

Target Protein
FGFR2

Protein Length
Partial

Protein Class
Druggable Genome, Protein Kinase, Secreted Protein, Transmembrane

Molecular Weight
39.5 kDa

TMD
1

Sequence
MVSWGRFICLVVVTMATLSLARPSFSLVEDTTLEPEEPPTKYQISQPEVYVAAPGESLEVRCLLKDAAVISWTKDGVHLGPNNRTVL

Product Description

Expression Systems
Sf9

Tag
C-DDK

Form
Powder

Purity

> 80% as determined by SDS-PAGE and Coomassie blue staining

Buffer

50mM Tris-HCl, pH8.0, 100mM glycine, 10% glycerol

Storage

Store at +4°C for up to one week or several months at -80°C

Target

Target Protein

FGFR2

Full Name

Fibroblast growth factor receptor 2

Introduction

The protein encoded by this gene is a member of the fibroblast growth factor receptor 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 is a high-affinity receptor for acidic, basic and/or keratinocyte growth factor, depending on the isoform. Mutations in this gene are associated with Crouzon syndrome, Pfeiffer syndrome, Craniosynostosis, Apert syndrome, Jackson-Weiss syndrome, Beare-Stevenson cutis gyrata syndrome, Saethre-Chotzen syndrome, and syndromic craniosynostosis. Multiple alternatively spliced transcript variants encoding different isoforms have been noted for this gene.

Alternative Names

BBDS; BEK; BFR-1; CD332; CEK3; CFD1; ECT1; JWS; K-SAM; KGFR; TK14; TK25; fibroblast growth factor receptor 2; fibroblast growth factor receptor; bacteria-expressed kinase; keratinocyte growth factor receptor; protein tyrosine kinase, receptor like 14; FGFR-2; K-sam

Gene ID

2263

UniProt ID

P21802