

# Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten! See the following pages for more information!



## Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

## Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

## SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

linkedin.com/company/szaboscandic in

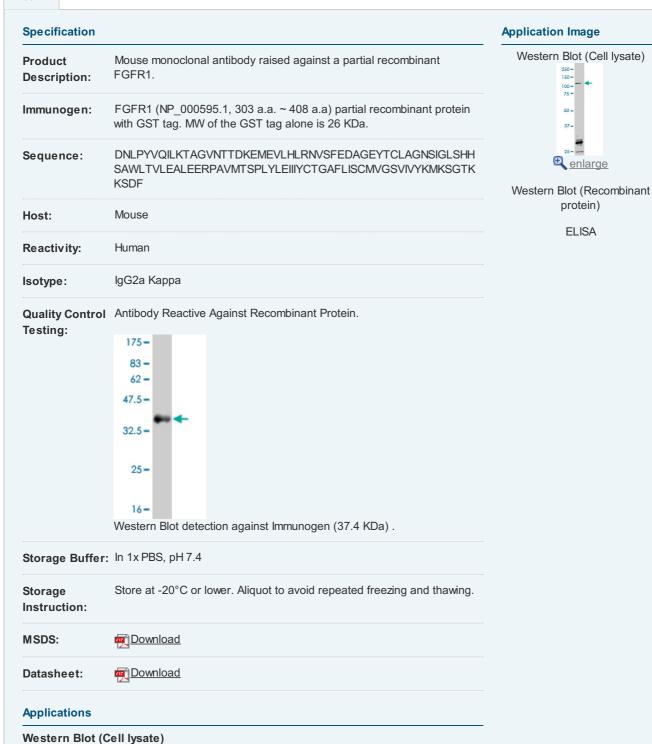




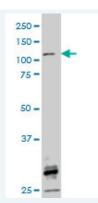
## FGFR1 monoclonal antibody (M15), clone 3C9

**Catalog #**: H00002260-M15 規格:[100 ug]

### List All



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FGFR1 monoclonal antibody (M15), clone 3C9 Western Blot analysis of FGFR1 expression in U-2 OS ( Cat # L022V1 ).

Protocol Download

Western Blot (Recombinant protein)

Protocol Download

**ELISA** 

Gene Information

Entrez GenelD: 2260

**GeneBank** 

NM 000604

Accession#:

Protein

NP 000595.1

Accession#:

Gene Name:

FGFR1

Gene Alias:

BFGFR,CD331,CEK,FGFBR,FLG,FLJ99988,FLT2,HBGFR,KAL2,N-

SAM

Gene

fibroblast growth factor receptor 1

**Description:** 

Omim ID:

101600, 123150, 136350, 147950

Gene Ontology: Hyperlink

Gene Summary: 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. [provided by RefSeq

Other

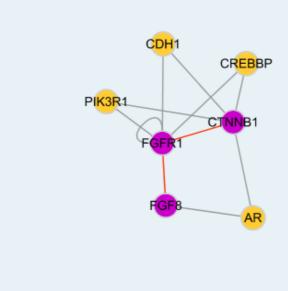
FMS-like tyrosine kinase

2,OTTHUMP00000190874,OTTHUMP00000190878,OTTHUMP0000019 Designations: 0879,OTTHUMP00000190881,basic fibroblast growth factor receptor

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1,fms-related tyrosine kinase 2,fms-related tyrosine kinase-2,heparinbinding growth factor receptor,hydroxyaryl

### Interactome



### **Gene Pathway**

Adherens junction MAPK signaling pathway Melanoma Pathways in cancer Prostate cancer Regulation of actin cytoskeleton

### **Related Disease**

Abnormalities, Multiple Acrocephalosyndactylia Alzheimer Disease Alzheimer disease Amenorrhea Anodontia Breast cancer Breast Neoplasms Bronchial Hyperreactivity Cardiovascular Diseases Chromosome Aberrations Chromosome Disorders Cleft Lip Cleft Palate Craniofacial Dysostosis Craniosynostoses Diabetes Complications Fractures, Bone Genetic Diseases, Inborn

... see more

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