# SZABO SCANDIC 

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Zuschläge

- Mindermengenzuschlag
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- Expressversand


## FGFR1(phospho Y154) \& FGFR1 Protein Phosphorylation Antibody Pair

Catalog \# : DP0034
規格: [1 Set]
List All


Application Image
In situ Proximity Ligation Assay (Cell)

| Storage | Store reagents of the antibody pair set at $-20^{\circ} \mathrm{C}$ or lower. Please aliquot |
| :---: | :---: |
| Ins | to avoid repeated freeze thaw cycle. Reagents should be returned to |
|  | $20^{\circ} \mathrm{C}$ storage immediately after use |

## Publication Reference

1. In situ detection of phosphorylated platelet-derived growth factor receptor beta using a generalized proximity ligation method. Jarvius M, Paulsson J, Weibrecht I, Leuchowius KJ, Andersson AC, Wahlby C, Gullberg M,Botling J, Sjoblom T, Markova B, Ostman A, Landegren U, Soderberg O. Mol Cell Proteomics. 2007 Sep;6(9):1500-9. Epub 2007 Jun 12.
2. Direct observation of individual endogenous protein complexes in situ by proximity ligation.
Soderberg O, Gullberg M, Jarvius M , Ridderstrale K, Leuchowius KJ, Jarvius J, Wester K, Hydbring P, Bahram F, Larsson LG, and Landegren U.
Nat Methods. 2006 Dec;3(12):995-1000. Epub 2006 Oct 29.
3. Cytokine detection by antibody-based proximity ligation.

Gullberg M, Gustafsdottir SM, Schallmeiner E, Jarvius J, Bjarnegard M, Betsholtz C, Landegren U, and Fredriksson S.
Proc Natl Acad Sci U S A. 2004 Jun 1;101(22):8420-4. Epub 2004 May 21.
4. Protein detection using proximity-dependent DNA ligation assays. Fredriksson S, GullbergM, Jarvius J, Olsson C, Pietras K, Gustafsdottir SM, Ostman A, and Landegren U .
Nat Biotechnol. 2002 May;20(5):473-7.
5. Highly specific detection of phosphorylated proteins by Duolink

Mats Gullberg and Ann-Catrin Andersson
Nature Methods 6. 2009

Applications
In situ Proximity Ligation Assay (Cell)

## Gene Information

Entrez GeneID: $\underline{2260}$
Gene Name: FGFR1

Gene Alias: BFGFR,CD331,CEK,FGFBR,FLG,FLJ99988,FLT2,HBGFR,KAL2,NSAM

Gene fibroblast growth factor receptor 1
Description:
Omim ID: $\quad 101600,123150,136350,147950$

## 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
Designations：2，OTTHUMP00000190874，OTTHUMP00000190878，OTTHUMP0000019 0879，OTTHUMP00000190881，basic fibroblast growth factor receptor 1，fms－related tyrosine kinase 2，fms－related tyrosine kinase－2，heparin－ binding 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

