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Lieferung & Zahlungsart

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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

SZABO-SCANDIC Handels GmbH

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](https://www.linkedin.com/company/szaboscandic)



ROR2 monoclonal antibody (M02), clone 1F7

Catalog # : H00004920-M02

規格 : [100 ug]

[List All](#)

Specification

Product Description: Mouse monoclonal antibody raised against a partial recombinant ROR2.

Immunogen: ROR2 (NP_004551, 34 a.a. ~ 143 a.a) partial recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.

Sequence: EVEVLDPNPLGLDGGDGIPTLKGYFLNFLEPVNNITVQGQTALHCK
VAGNPPPNVRLKNDAPVWQEPRIIRKTEYGSRLRIQDLDTTDTGYQ
CVATNGMKT

Host: Mouse

Reactivity: Human

Isotype: IgG2a Kappa

Quality Control Testing: Antibody Reactive Against Recombinant Protein.

Storage Buffer: In 1x PBS, pH 7.4

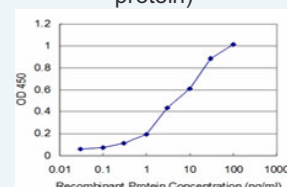
Storage Instruction: Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

MSDS:  [Download](#)

Datasheet:  [Download](#)

Application Image

Sandwich ELISA (Recombinant protein)

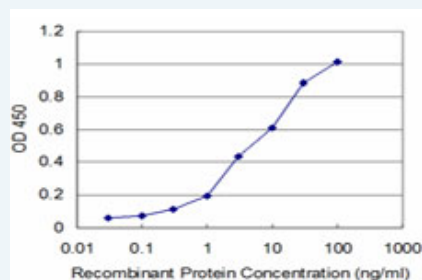


 [enlarge](#)

ELISA

Applications

Sandwich ELISA (Recombinant protein)



Detection limit for recombinant GST tagged ROR2 is approximately 0.3ng/ml as a capture antibody.

 [Protocol Download](#)

ELISA

Gene Information

Entrez GeneID: [4920](#)

GeneBank [NM_004560](#)

Accession#:**Protein** [NP_004551](#)**Accession#:****Gene Name:** ROR2**Gene Alias:** BDB,BDB1,MGC163394,NTRKR2**Gene** receptor tyrosine kinase-like orphan receptor 2**Description:****Omim ID:** [113000](#), [268310](#), [602337](#)**Gene Ontology:** [Hyperlink](#)

Gene Summary: The protein encoded by this gene is a receptor protein tyrosine kinase and type I transmembrane protein that belongs to the ROR subfamily of cell surface receptors. The protein may be involved in the early formation of the chondrocytes and may be required for cartilage and growth plate development. Mutations in this gene can cause brachydactyly type B, a skeletal disorder characterized by hypoplasia/aplasia of distal phalanges and nails. In addition, mutations in this gene can cause the autosomal recessive form of Robinow syndrome, which is characterized by skeletal dysplasia with generalized limb bone shortening, segmental defects of the spine, brachydactyly, and a dysmorphic facial appearance. [provided by RefSeq]

Other OTTHUMP00000021634,OTTHUMP00000063680,neurotrophic tyrosine
Designations: kinase receptor-related 2,tyrosine-protein kinase transmembrane receptor ROR2

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