

ROR2 Antibody (C-term)
Purified Rabbit Polyclonal Antibody (Pab)
Catalog # AP7672b

Specification

ROR2 Antibody (C-term) - Product Information

Application	WB, IHC-P,E
Primary Accession	Q01974
Reactivity	Human
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit Ig
Antigen Region	915-943

ROR2 Antibody (C-term) - Additional Information

Gene ID 4920

Other Names

Tyrosine-protein kinase transmembrane receptor ROR2, Neurotrophic tyrosine kinase, receptor-related 2, ROR2, NTRKR2

Target/Specificity

This ROR2 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 915-943 amino acids from the C-terminal region of human ROR2.

Dilution

WB~~1:1000
IHC-P~~1:50~100

Format

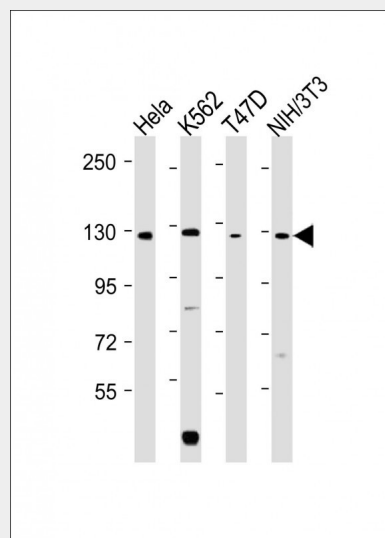
Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is prepared by Saturated Ammonium Sulfate (SAS) precipitation followed by dialysis against PBS.

Storage

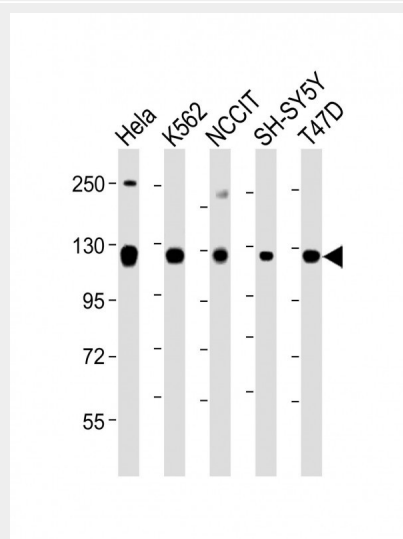
Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

ROR2 Antibody (C-term) is for research use only and not for use in diagnostic or therapeutic procedures.



All lanes : Anti-ROR2 Antibody (C-term) at 1:2000 dilution Lane 1: HeLa whole cell lysate Lane 2: K562 whole cell lysate Lane 3: T47D whole cell lysate Lane 4: NIH/3T3 whole cell lysate Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution. Predicted band size : 105 kDa Blocking/Dilution buffer: 5% NFDM/TBST.



All lanes : Anti-ROR2 Antibody (C-term) at 1:1000-1:2000 dilution Lane 1: HeLa whole

ROR2 Antibody (C-term) - Protein Information

Name ROR2

Synonyms NTRKR2

Function

Tyrosine-protein kinase receptor which may be involved in the early formation of the chondrocytes. It seems to be required for cartilage and growth plate development (By similarity). Phosphorylates YWHAB, leading to induction of osteogenesis and bone formation (PubMed:17717073). In contrast, has also been shown to have very little tyrosine kinase activity in vitro. May act as a receptor for wnt ligand WNT5A which may result in the inhibition of WNT3A-mediated signaling (PubMed:25029443).

Cellular Location

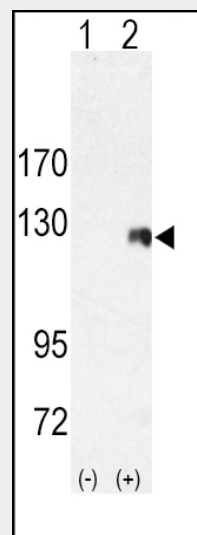
Cell membrane; Single-pass type I membrane protein

ROR2 Antibody (C-term) - Protocols

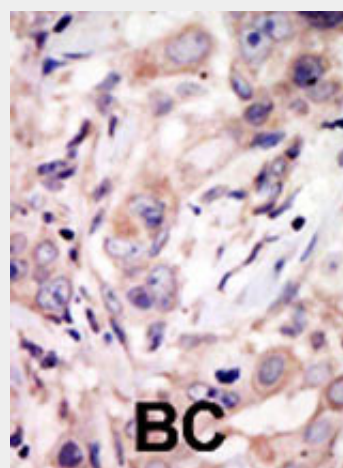
Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

cell lysate Lane 2: K562 whole cell lysate
Lane 3: NCCIT whole cell lysate Lane 4:
SH-SY5Y whole cell lysate Lane 5: T47D
whole cell lysate Lysates/proteins at 20 µg
per lane. Secondary Goat Anti-Rabbit IgG,
(H+L), Peroxidase conjugated at 1/10000
dilution. Predicted band size : 105 kDa
Blocking/Dilution buffer: 5% NFDM/TBST.



Western blot analysis of ROR2 (arrow) using rabbit polyclonal ROR2 Antibody (C-term) (Cat.#AP7672b). 293 cell lysates (2 ug/lane) either nontransfected (Lane 1) or transiently transfected with the ROR2 gene (Lane 2) (Origene Technologies).



Formalin-fixed and paraffin-embedded human cancer tissue reacted with the primary antibody, which was peroxidase-conjugated to the secondary antibody, followed by DAB staining. This data demonstrates the use of this antibody for immunohistochemistry; clinical relevance has not been evaluated. BC = breast carcinoma;

HC = hepatocarcinoma.

ROR2 Antibody (C-term) - Background

ROR2 is a tyrosine-protein kinase receptor which may be involved in the early formation of the chondrocytes. It seems to be required for cartilage and growth plate development. This Type I membrane protein is expressed at high levels during early embryonic development. The expression levels drop strongly around day 16 and there are only very low levels in adult tissues. Defects in ROR2 are a cause of brachydactyly type B1 (BDB1). BDB1 is an autosomal dominant skeletal disorder characterized by hypoplasia/aplasia of distal phalanges and nails. In BDB1 the middle phalanges are short but in addition the terminal phalanges are rudimentary or absent. Both fingers and toes are affected. The thumbs and big toes are usually deformed. Defects in ROR2 are a cause of recessive Robinow syndrome (RRS). RRS is an autosomal disorder characterized by skeletal dysplasia with generalized limb bone shortening, segmental defects of the spine, brachydactyly and a dysmorphic facial appearance. The protein contains 1 frizzled (FZ) domain, 1 immunoglobulin-like C2-type domain, and 1 kringle domain.

ROR2 Antibody (C-term) - References

Afzal, A.R., et al., Nat. Genet. 25(4):419-422 (2000).
Oldridge, M., et al., Nat. Genet. 24(3):275-278 (2000).
van Bokhoven, H., et al., Nat. Genet. 25(4):423-426 (2000).
Schwabe, G.C., et al., Am. J. Hum. Genet. 67(4):822-831 (2000).
Masiakowski, P., et al., J. Biol. Chem. 267(36):26181-26190 (1992).