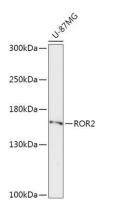
## **ROR2 Rabbit Polyclonal Antibody**

## CAB5620



Product Information	Protein Background
Size:	The protein encoded by this gene is a receptor protein tyrosine kinase and type
20uL, 50uL, 100uL, 200uL	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,
Observed MW:	
135kDa	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
Calculated MW:	of the spine, brachydactyly, and a dysmorphic facial appearance.
104kDa	Immunogen information
Applications:	Gene ID:
WB IHC	4920
Reactivity:	Uniprot Q01974
Human, Mouse, Rat	
	Synonyms:
Antibody Information	ROR2; BDB; BDB1; NTRKR2
<b>Recommended dilutions:</b> WB 1:500 - 1:2000 IHC 1:100 - 1:200	Immunogen:
Source: Rabbit	A synthetic peptide corresponding to a sequence within amino acids 100-200 of human ROR2 (NP_004551.2).
<b>lsotype:</b> lgG	<b>Storage:</b> Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

**Purification:** Affinity purification



Western blot analysis of extracts of U-87MG cells, using ROR2 antibody (CAB5620) at 1:1000 dilution. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (CABS014) at 1:10000 dilution. Lysates/proteins: 25ug per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (CABM00020). Exposure time: 3MIN.