

Product datasheet for AP54739PU-N

ZRANB1 (C-term) Rabbit Polyclonal Antibody

Product data:

Product Type: Primary Antibodies

Applications: IHC, WB

Recommended Dilution: Peptide ELISA: 1/1000.

Western blot: 1/100~1/500.

Immunohistochemistry: 1/50~1/100.

Reactivity: Human, Mouse

Host: Rabbit

Isotype: lg

Clonality: Polyclonal

Immunogen: KLH conjugated synthetic peptide between 665~692 amino acids from the C-terminal region

of human ZRANB1

Specificity: This antibody detects ZRANB1 (C-term).

Formulation: PBS with 0.09% (W/V) sodium azide

State: Aff - Purified

State: Liquid Ig fraction

Concentration: lot specific

Purification: Protein A column followed by peptide affinity purification

Conjugation: Unconjugated

Store undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer. Storage:

Avoid repeated freezing and thawing.

Stability: Shelf life: one year from despatch.

Gene Name: Homo sapiens zinc finger RANBP2-type containing 1 (ZRANB1)

Database Link: Entrez Gene 360216 MouseEntrez Gene 54764 Human

Q9UGI0

Background: ZRANB1 is a positive regulator of the Wnt signaling pathway that specifically cleaves 'Lys-63'-

linked ubiquitin chains. It acts by deubiquitinating APC protein, a negative regulator of Wnt-

mediated transcription and may also modulate TNF-alpha signaling.

Synonyms: **TRABID**



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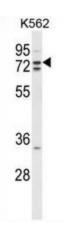
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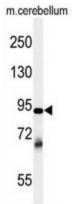
Note: Molecular Weight: 80967 Da

Protein Families: Druggable Genome

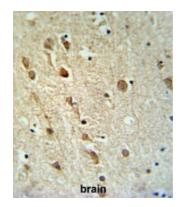
Product images:



Western blot analysis of ZRANB1 Antibody (Cterm) in K562 cell line lysates (35 ug/lane). ZRANB1 (arrow) was detected using the purified Pab.



Western blot analysis of ZRANB1 Antibody (Cterm) in mouse cerebellum tissue lysates (35 ug/lane). ZRANB1 (arrow) was detected using the purified Pab.



ZRANB1 Antibody (C-term) IHC analysis in formalin fixed and paraffin embedded brain tissue followed by peroxidase conjugation of the secondary antibody and DAB staining. This data demonstrates the use of the ZRANB1 Antibody (C-term) for immunohistochemistry. Clinical relevance has not been evaluated.