

**RanGAP1 (Ran-GTPase) Antibody (Sumoylation Site Specific) Blocking peptide**  
**Synthetic peptide**  
**Catalog # BP2503a****Specification**

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**RanGAP1 (Ran-GTPase) Antibody (Sumoylation Site Specific) Blocking peptide - Product Information**Primary Accession  
Other Accession[P46060](#)  
[NP\\_002874](#)**RanGAP1 (Ran-GTPase) Antibody (Sumoylation Site Specific) Blocking peptide - Additional Information****Gene ID** 5905**Other Names**

Ran GTPase-activating protein 1, RanGAP1, RANGAP1, KIAA1835, SD

**Target/Specificity**

The synthetic peptide sequence used to generate the antibody [AP2503a](/product/products/AP2503a) was selected from the region of human Ran-GTPase Sumoylation site. A 10 to 100 fold molar excess to antibody is recommended. Precise conditions should be optimized for a particular assay.

**Format**

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

**Storage**

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

**Precautions**

This product is for research use only. Not for use in diagnostic or therapeutic procedures.

**RanGAP1 (Ran-GTPase) Antibody (Sumoylation Site Specific) Blocking peptide - Protein Information****Name** RANGAP1**Synonyms** KIAA1835, SD**Function**

GTPase activator for RAN (PubMed: [8146159](http://www.uniprot.org/citations/8146159), PubMed: [8896452](http://www.uniprot.org/citations/8896452), PubMed: [16428860](http://www.uniprot.org/citations/16428860)). Converts cytoplasmic GTP-bound RAN to GDP-bound RAN, which is essential for RAN-mediated nuclear import and export (PubMed: [8896452](http://www.uniprot.org/citations/8896452), PubMed: [8896452](http://www.uniprot.org/citations/8896452)).

href="http://www.uniprot.org/citations/27160050" target="\_blank">27160050</a>). Mediates dissociation of cargo from nuclear export complexes containing XPO1, RAN and RANBP2 after nuclear export (PubMed:<a href="http://www.uniprot.org/citations/27160050" target="\_blank">27160050</a>).

#### **Cellular Location**

Cytoplasm. Nucleus, nucleoplasm. Nucleus envelope. Chromosome, centromere, kinetochore. Cytoplasm, cytoskeleton, spindle. Note=Cytoplasmic during interphase Detected at the nuclear envelope during interphase (PubMed:11854305, PubMed:15037602). Targeted to the nuclear pores after sumoylation (PubMed:11854305). During mitosis, associates with mitotic spindles, but is essentially not detected at the spindle poles (PubMed:11854305, PubMed:15037602). Association with kinetochores appears soon after nuclear envelope breakdown and persists until late anaphase (PubMed:11854305). Mitotic location also requires sumoylation (PubMed:11854305).

#### **Tissue Location**

Highly expressed in brain, thymus and testis.

### **RanGAP1 (Ran-GTPase) Antibody (Sumoylation Site Specific) Blocking peptide - Protocols**

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

- [Blocking Peptides](#)

### **RanGAP1 (Ran-GTPase) Antibody (Sumoylation Site Specific) Blocking peptide - Images**

### **RanGAP1 (Ran-GTPase) Antibody (Sumoylation Site Specific) Blocking peptide - Background**

RanGAP1, is a homodimeric 65-kD polypeptide that specifically induces the GTPase activity of RAN, but not of RAS by over 1,000-fold. RanGAP1 is the immediate antagonist of RCC1, a regulator molecule that keeps RAN in the active, GTP-bound state. The RANGAP1 gene encodes a 587-amino acid polypeptide. The sequence is unrelated to that of GTPase activators for other RAS-related proteins, but is 88% identical to Fug1, the murine homolog of yeast Rna1p. RanGAP1 and RCC1 control RAN-dependent transport between the nucleus and cytoplasm. RanGAP1 is a key regulator of the RAN GTP/GDP cycle.

### **RanGAP1 (Ran-GTPase) Antibody (Sumoylation Site Specific) Blocking peptide - References**

Bischoff, F.R., et al., Proc. Natl. Acad. Sci. U.S.A. 92(5):1749-1753 (1995). Bischoff, F.R., et al., Proc. Natl. Acad. Sci. U.S.A. 91(7):2587-2591 (1994). Matunis, M.J., et al., J. Cell Biol. 135 (6 Pt 1), 1457-1470 (1996).