

**Aurora B Rabbit mAb**  
Catalog # AP75131**Specification**

---

**Aurora B Rabbit mAb - Product Information**

Application	<b>WB</b>
Primary Accession	<a href="#">O96GD4</a>
Reactivity	<b>Human</b>
Host	<b>Rabbit</b>
Clonality	<b>Monoclonal Antibody</b>
Calculated MW	<b>39311</b>

**Aurora B Rabbit mAb - Additional Information****Gene ID** 9212**Other Names**

AURKB

**Dilution**

WB~~1/500-1/1000

**Format**

Liquid

**Aurora B Rabbit mAb - Protein Information****Name** AURKB**Function**

Serine/threonine-protein kinase component of the chromosomal passenger complex (CPC), a complex that acts as a key regulator of mitosis (PubMed: [11516652](http://www.uniprot.org/citations/11516652), PubMed: [12925766](http://www.uniprot.org/citations/12925766), PubMed: [14610074](http://www.uniprot.org/citations/14610074), PubMed: [14722118](http://www.uniprot.org/citations/14722118), PubMed: [29449677](http://www.uniprot.org/citations/29449677)). The CPC complex has essential functions at the centromere in ensuring correct chromosome alignment and segregation and is required for chromatin-induced microtubule stabilization and spindle assembly (PubMed: [11516652](http://www.uniprot.org/citations/11516652), PubMed: [12925766](http://www.uniprot.org/citations/12925766), PubMed: [14610074](http://www.uniprot.org/citations/14610074), PubMed: [14722118](http://www.uniprot.org/citations/14722118), PubMed: [26829474](http://www.uniprot.org/citations/26829474)). Involved in the bipolar attachment of spindle microtubules to kinetochores and is a key regulator for the onset of cytokinesis during mitosis (PubMed: [15249581](http://www.uniprot.org/citations/15249581)). Required for central/midzone spindle assembly and cleavage furrow formation (PubMed: [15249581](http://www.uniprot.org/citations/15249581)).

href="http://www.uniprot.org/citations/12458200" target="\_blank">12458200</a>, PubMed:<a href="http://www.uniprot.org/citations/12686604" target="\_blank">12686604</a>). Key component of the cytokinesis checkpoint, a process required to delay abscission to prevent both premature resolution of intercellular chromosome bridges and accumulation of DNA damage: phosphorylates CHMP4C, leading to retain abscission-competent VPS4 (VPS4A and/or VPS4B) at the midbody ring until abscission checkpoint signaling is terminated at late cytokinesis (PubMed:<a href="http://www.uniprot.org/citations/22422861" target="\_blank">22422861</a>, PubMed:<a href="http://www.uniprot.org/citations/24814515" target="\_blank">24814515</a>). AURKB phosphorylates the CPC complex subunits BIRC5/survivin, CDCA8/borealin and INCENP (PubMed:<a href="http://www.uniprot.org/citations/11516652" target="\_blank">11516652</a>, PubMed:<a href="http://www.uniprot.org/citations/12925766" target="\_blank">12925766</a>, PubMed:<a href="http://www.uniprot.org/citations/14610074" target="\_blank">14610074</a>). Phosphorylation of INCENP leads to increased AURKB activity (PubMed:<a href="http://www.uniprot.org/citations/11516652" target="\_blank">11516652</a>, PubMed:<a href="http://www.uniprot.org/citations/12925766" target="\_blank">12925766</a>, PubMed:<a href="http://www.uniprot.org/citations/14610074" target="\_blank">14610074</a>). Other known AURKB substrates involved in centromeric functions and mitosis are CENPA, DES/desmin, GPAF, KIF2C, NSUN2, RACGAP1, SEPTIN1, VIM/vimentin, HASPIN, and histone H3 (PubMed:<a href="http://www.uniprot.org/citations/11756469" target="\_blank">11756469</a>, PubMed:<a href="http://www.uniprot.org/citations/11784863" target="\_blank">11784863</a>, PubMed:<a href="http://www.uniprot.org/citations/11856369" target="\_blank">11856369</a>, PubMed:<a href="http://www.uniprot.org/citations/12689593" target="\_blank">12689593</a>, PubMed:<a href="http://www.uniprot.org/citations/14602875" target="\_blank">14602875</a>, PubMed:<a href="http://www.uniprot.org/citations/16103226" target="\_blank">16103226</a>, PubMed:<a href="http://www.uniprot.org/citations/21658950" target="\_blank">21658950</a>). A positive feedback loop involving HASPIN and AURKB contributes to localization of CPC to centromeres (PubMed:<a href="http://www.uniprot.org/citations/21658950" target="\_blank">21658950</a>). Phosphorylation of VIM controls vimentin filament segregation in cytokinetic process, whereas histone H3 is phosphorylated at 'Ser-10' and 'Ser-28' during mitosis (H3S10ph and H3S28ph, respectively) (PubMed:<a href="http://www.uniprot.org/citations/11784863" target="\_blank">11784863</a>, PubMed:<a href="http://www.uniprot.org/citations/11856369" target="\_blank">11856369</a>). AURKB is also required for kinetochore localization of BUB1 and SGO1 (PubMed:<a href="http://www.uniprot.org/citations/15020684" target="\_blank">15020684</a>, PubMed:<a href="http://www.uniprot.org/citations/17617734" target="\_blank">17617734</a>). Phosphorylation of p53/TP53 negatively regulates its transcriptional activity (PubMed:<a href="http://www.uniprot.org/citations/20959462" target="\_blank">20959462</a>). Key regulator of active promoters in resting B- and T-lymphocytes: acts by mediating phosphorylation of H3S28ph at active promoters in resting B-cells, inhibiting RNF2/RING1B-mediated ubiquitination of histone H2A and enhancing binding and activity of the USP16 deubiquitinase at transcribed genes (By similarity). Acts as an inhibitor of CGAS during mitosis: catalyzes phosphorylation of the N-terminus of CGAS during the G2-M transition, blocking CGAS liquid phase separation and activation, and thereby preventing CGAS-induced autoimmunity (PubMed:<a href="http://www.uniprot.org/citations/33542149" target="\_blank">33542149</a>). Phosphorylates KRT5 during anaphase and telophase (By similarity). Phosphorylates ATXN10 which promotes phosphorylation of ATXN10 by PLK1 and may play a role in the regulation of cytokinesis and stimulating the proteasomal degradation of ATXN10 (PubMed:<a href="http://www.uniprot.org/citations/25666058" target="\_blank">25666058</a>).

### Cellular Location

Nucleus. Chromosome. Chromosome, centromere. Chromosome, centromere, kinetochore. Cytoplasm, cytoskeleton, spindle. Midbody. Note=Localizes on chromosome arms and inner centromeres from prophase through metaphase and then transferring to the spindle midzone and midbody from anaphase through cytokinesis (PubMed:20929775). Colocalized with gamma tubulin in the midbody (PubMed:17726514). Proper localization of the active, Thr-232- phosphorylated form during metaphase may be dependent upon interaction with SPDYC (PubMed:20605920). Colocalized with SIRT2 during cytokinesis with the midbody (PubMed:17726514). Localization (and probably targeting of the CPC) to the inner centromere occurs predominantly in regions with

overlapping mitosis-specific histone phosphorylations H3pT3 and H2ApT12 (PubMed:20929775).

#### Tissue Location

High level expression seen in the thymus. It is also expressed in the spleen, lung, testis, colon, placenta and fetal liver. Expressed during S and G2/M phase and expression is up-regulated in cancer cells during M phase.

#### Aurora B Rabbit mAb - 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](#)

#### Aurora B Rabbit mAb - Images

