

Aurora B Antibody

Rabbit mAb

Catalog # AP90074

Product Information

Application	WB, IHC, IF, ICC, IP, IHF
Primary Accession	Q96GD4
Reactivity	Human
Clonality	Monoclonal
Other Names	AIK2; AIM1; ARK2; AURKB; Aurora- and Ipl1-like midbody-associated protein 1; Aurora-B; Aurora/IPL1-related kinase 2; STK-1; STK12;
Isotype	Rabbit IgG
Host	Rabbit
Calculated MW	39311

Additional Information

Dilution	WB 1:1000~1:2000 IHC 1:50~1:200 ICC/IF 1:50~1:200 IP 1:50
Purification	Affinity-chromatography
Immunogen	A synthesized peptide derived from human Aurora B
Description	May be directly involved in regulating the cleavage of polar spindle microtubules and is a key regulator for the onset of cytokinesis during mitosis. Component of the chromosomal passenger complex (CPC), a complex that acts as a key regulator of mitosis. 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.
Storage Condition and Buffer	Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at +4°C short term. Store at -20°C long term. Avoid freeze / thaw cycle.

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 , PubMed: 12925766 , PubMed: 14610074 , PubMed: 14722118 , PubMed: 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 , PubMed: 12925766 , PubMed: 14610074 , PubMed: 14722118 , PubMed: 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). Required for central/midzone spindle assembly and cleavage furrow

formation (PubMed:[12458200](#), PubMed:[12686604](#)). 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:[22422861](#), PubMed:[24814515](#)). AURKB phosphorylates the CPC complex subunits BIRC5/survivin, CDCA8/borealin and INCENP (PubMed:[11516652](#), PubMed:[12925766](#), PubMed:[14610074](#)). Phosphorylation of INCENP leads to increased AURKB activity (PubMed:[11516652](#), PubMed:[12925766](#), PubMed:[14610074](#)). 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:[11756469](#), PubMed:[11784863](#), PubMed:[11856369](#), PubMed:[12689593](#), PubMed:[14602875](#), PubMed:[16103226](#), PubMed:[21658950](#)). A positive feedback loop involving HASPIN and AURKB contributes to localization of CPC to centromeres (PubMed:[21658950](#)). 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:[11784863](#), PubMed:[11856369](#)). AURKB is also required for kinetochore localization of BUB1 and SGO1 (PubMed:[15020684](#), PubMed:[17617734](#)). Phosphorylation of p53/TP53 negatively regulates its transcriptional activity (PubMed:[20959462](#)). 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:[33542149](#)). 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:[25666058](#)).

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.

Images

Western blot analysis of Aurora B expression in HeLa cell lysate.

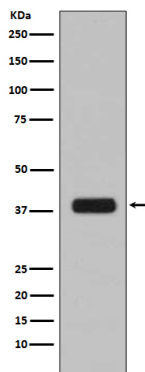


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Immunohistochemical analysis of paraffin-embedded human tonsil, using Aurora B Antibody.

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