

Datasheet for ABIN968357

anti-Aurora Kinase B antibody (AA 2-124)





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Overview	
Quantity:	50 µg
Target:	Aurora Kinase B (AURKB)
Binding Specificity:	AA 2-124
Reactivity:	Human, Mouse, Rat
Host:	Mouse
Clonality:	Monoclonal
Conjugate:	This Aurora Kinase B antibody is un-conjugated
Application:	Western Blotting (WB), Immunofluorescence (IF)
Product Details	
Immunogen:	Rat AIM-1 aa. 2-124
Clone:	6-AIM
Isotype:	lgG1
Cross-Reactivity:	Human, Mouse (Murine)
Characteristics:	1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
	2. Source of all serum proteins is from USDA inspected abattoirs located in the United States.
	3. Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide
	compounds in running water before discarding to avoid accumulation of potentially explosive

Purification: The monoclonal

deposits in plumbing.

4. Please refer to us for technical protocols.

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity

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Target Details

Target:	Aurora Kinase B (AURKB)
Alternative Name:	AIM-1 (AURKB Products)
Background:	The mitotic phase of the cell cycle is a complex process that ensures the fidelity of
	chromosome segregation. During the final stage of mitosis (telophase), segregated
	chromosomes become less condense and nuclear membranes surround the two sets of
	daughter chromosomes. Simultaneously, the separation and segregation of the cytoplasm
	(cytokinesis) ensures complete division and formation of two identical daughter cells.
	Regulation of cytokinesis is poorly understood and errors in this process can lead to cell death
	or oncogenesis. The Drosophila serine/threonine protein kinase Aurora and the S. cerevisiae
	Ipl1 kinase are highly homologous and are required for progression through mitosis. Their
	mammalian homolog AIM-1 (also known as Aurora and IpI1-like midbody associated protein)
	accumulates at the G2/M interface. During late anaphase, AIM-1 is found at the equator of
	central spindles. However, during telophase and cytokinesis, it is found at the midbody.
	Although over-expression of a kinase-inactive AIM-1 mutant disrupts formation of the cleavage
	furrow, nuclear division is unaffected. Thus, it is thought that AIM-1 is essential for cleavage
	furrowing and the onset of cytokinesis.
	Synonyms: Aurora B, Aurora and Ipl1-like midbody associated protein
Molecular Weight:	41 kDa
Pathways:	TCR Signaling, Cell Division Cycle, Maintenance of Protein Location, Hepatitis C, Toll-Like
	Receptors Cascades

Application Details

Comment:	Related Products: ABIN967389, ABIN968537
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Format: Concentration:	Liquid 250 μg/mL

Handling

Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C
Storage Comment:	Store undiluted at -20°C.

Publications

Product cited in:

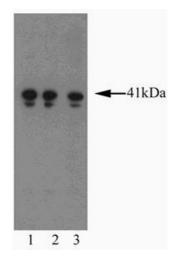
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Lange, Rebollo, Herold, González: "Cdc37 is essential for chromosome segregation and cytokinesis in higher eukaryotes." in: **The EMBO journal**, Vol. 21, Issue 20, pp. 5364-74, (2002) (PubMed).

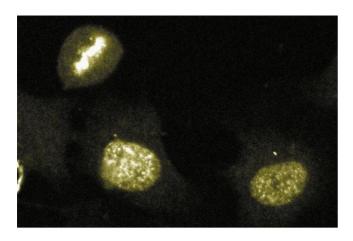
Tatsuka, Katayama, Ota, Tanaka, Odashima, Suzuki, Terada: "Multinuclearity and increased ploidy caused by overexpression of the aurora- and IpI1-like midbody-associated protein mitotic kinase in human cancer cells." in: **Cancer research**, Vol. 58, Issue 21, pp. 4811-6, (1998) (PubMed).

Terada, Tatsuka, Suzuki, Yasuda, Fujita, Otsu: "AIM-1: a mammalian midbody-associated protein required for cytokinesis." in: **The EMBO journal**, Vol. 17, Issue 3, pp. 667-76, (1998) (PubMed).



Western Blotting

Image 1. Western blot analysis of AIM-1 on a Jurkat cell lysate (Human T-cell leukemia, ATCC TIB-152). Lane 1: 1:250, lane 2: 1:500, lane 3: 1:1000 dilution of the Mouse Anti-AIM-1 antibody.



Immunofluorescence

Image 2. Immunofluorescence staining of human endothelial cells.