BIOMOL GmbH Waidmannstr. 35 biomol 22769 Hamburg info@biomol.de www.biomol.de Phone: +49-40-8532600 or 0800-2466651 (D) Fax: +49-40-85326022 or 0800-2466652 (D)



Mouse Monoclonal Antibody to

aurora A N-Terminus

clone 7F12

Order No.:	0233-100/auroraA-7F12			
Size (ug)	100			

Size (µg) Lot No.:

0233S



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Isotype	Species Reactivity	Applications	Mol. Weight	Ref.Cell Line	Epitope		Immunogen
lgG1	human, mouse, rat, dog	WB	47 kDa	A431	N-terminus		peptide conjugated to hemocyanin
Backgrour	nd and Specificity:					Related Pro	<u>ducts</u>

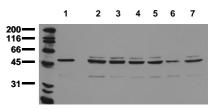
Aurora proteins are members of a serine/threonine kinase family. They play a crucial role in mitosis by regulating chromosome segregation and cytokinesis. There are three forms of Aurora proteins in mammalian cells: AuroraA, B and C. AuroraA (Aurora-2; STK6, ARK1, Aurora/IPL-1 related kinase) associates with centrosomes and microtubules during mitosis. Phosphorylation of a threonine residue within the activation loop of the catalytic domain lead to activation of AuroraA. AuroraB (Aurora-1) is responsible for chromatin modification and histone H3 phosphporylation.

mab to aurora A/B #0245-100/auroraA/B-5F11

01/1603075

Purification:	The antibody was purified from serum-free cell culture supernatant by subsequent ultrafiltration and size exclusion chromatography.
Formulation:	lyophilized from 1 ml PBS / 0.09 % Na-azide / PEG and Sucrose
Reconstitution:	Reconstitute with 1 ml H_2O (15 min, RT).
Stability:	For long-term storage, freeze lyophilizate upon arrival (-20°C). Upon reconstitution, aliquote and freeze in liquid nitrogen; reconstituted antibody can be stored frozen at -80°C up to 1 year. Thaw aliquots at 37°C. Thawed aliquots may be stored at 4°C up to 3 months.
	Avoid repeated freeze / thaw cycles.
Positive Control:	#0831: Cell Lysate from untreated A431 cells.
Immunoblotting:	0.5 μg/ml for HRPO/ECL detection <u>Recommended blocking buffer:</u> Casein/Tween 20 based blocking and blot incubation buffer, e.g. nanoTools product #3031-500/CPPT or #3031-3000/CPPT.
Immunoprecipitation:	ND
Immunocytochemistry:	ND
ELISA:	ND

All products are supplied for research and investigational use only. Not for use in humans or laboratory animals.



Detection of endogenous auroraA

Whole cell extracts of vanadate treated tumor cells (20.000 cells per lane) were applied to SDS-PAGE and transferred to a PVDF membrane. The immunoblot was probed with mab auroraA-7F12 (0.5 $\mu\text{g}/$ ml) for 1h at RT and developed by ECL (exp. time: 30 sec)

lane 1: A431; lane 2: SW480; lane 3: SW620; lane 4: HT29; lane 5: MCF-7; lane 6: MDA-MB231; lane 7: T47D