

# Product Datasheet

## Aurora A [p Thr288] Antibody - BSA Free NBP3-05434

Unit Size: 100 ul

Store at -20C. Avoid freeze-thaw cycles.

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**NBP3-05434**

Aurora A [p Thr288] Antibody - BSA Free

Product Information	
Unit Size	100 ul
Concentration	Please see the vial label for concentration. If unlisted please contact technical services.
Storage	Store at -20C. Avoid freeze-thaw cycles.
Clonality	Polyclonal
Preservative	0.02% Sodium Azide
Isotype	IgG
Purity	Affinity purified
Buffer	PBS (pH 7.3), 50% glycerol
Target Molecular Weight	46 kDa

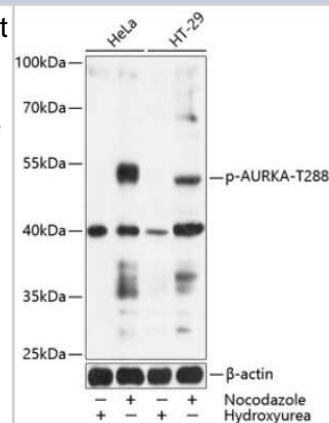
Product Description	
Description	Novus Biologicals Rabbit Aurora A [p Thr288] Antibody - BSA Free (NBP3-05434) is a polyclonal antibody validated for use in WB, ELISA and Simple Western. Anti-Aurora A Antibody: Cited in 1 publication. All Novus Biologicals antibodies are covered by our 100% guarantee.
Host	Rabbit
Gene ID	6790
Gene Symbol	AURKA
Species	Human
Immunogen	A synthetic phosphorylated peptide around T288 of human Aurora A (NP_003591.2). RTTLC

Product Application Details	
Applications	Western Blot, Simple Western, ELISA
Recommended Dilutions	Western Blot 1:500-1:2000, Simple Western, ELISA
Application Notes	See <a href="#">Simple Western Antibody Database</a> for Simple Western validation: Tested in CHLA-10, CHLA-258, CHLA-32, CHLA-9, COG-E-352, RD-ES, SK-ES-1, TC-32, TC-71, separated by Size

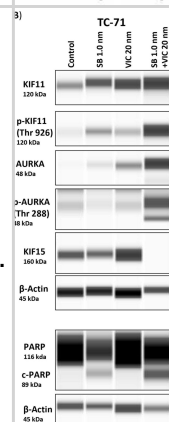


## Images

Western Blot: Aurora A [p Thr288] Antibody [NBP3-05434] - Western blot analysis of extracts of HeLa and HT-29 cells, using Aurora A antibody (NBP3-05434) at 1:1000 dilution. HeLa cells were treated by Hydroxyurea (4mM) for 20 hours. HeLa cells were treated by Nocodazole (50ng/mL) for 20 hours. HT-29 cells were treated by Hydroxyurea (4 mM) for 20 hours or treated by Nocodazole (100ng/mL) for 16 hours. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) at 1:10000 dilution. Lysates/proteins: 25ug per lane. Blocking buffer: 3% BSA. Detection: ECL Basic Kit. Exposure time: 90s.



Simple Western: Aurora A [p Thr288] Antibody - BSA Free [NBP3-05434] - Analysis of protein expression post-drug treatment. (A) CHLA-10 and (B) TC-71 cells treated with drugs were assessed for changes in protein expression 24 h post-treatment via capillary electrophoresis-based Wes analysis. Increased protein levels of KIF11, p-KIF11Thr926 AURKA, and p-AURKAThr288 were observed for the drug combination group, whereas KIF15 levels were noticeably lower. Similarly, enhanced cleaved-PARP expression was observed with the combination treatment. The uncropped blots are shown in Figures S8 and S9. Image collected and cropped by CiteAb from the following publication (<https://pubmed.ncbi.nlm.nih.gov/37894278>), licensed under a CC-BY license. Not internally tested by Novus Biologicals.



## Publications

Turaga SM, Vishwakarma V, Hembruff SL et al. Inducing Mitotic Catastrophe as a Therapeutic Approach to Improve Outcomes in Ewing Sarcoma Cancers 2023-10-10 [PMID: 37894278] (Simple Western, Human)



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### **Products Related to NBP3-05434**

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NBP2-33376H	Blue Marker Antibody (6F4-F6) [HRP]
HAF008	Goat anti-Rabbit IgG Secondary Antibody [HRP]
NB7160	Goat anti-Rabbit IgG (H+L) Secondary Antibody [HRP]
NBP2-24891	Rabbit IgG Isotype Control

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### **Limitations**

This product is for research use only and is not approved for use in humans or in clinical diagnosis. Primary Antibodies are guaranteed for 1 year from date of receipt.

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