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**Research Use Only. Not for
diagnostic or therapeutic use.**

EB05035 - Goat Anti-Aurora kinase A / AURKA Antibody

Size: 100µg specific antibody in 200µl



Target Protein

Principal Names: AURKA, aurora kinase A, AIK, ARK1, AURA, AURORA2, BTAK, MGC34538, STK15, STK6, STK7, IPL1-related kinase, aurora-A, aurora-related kinase 1, aurora/IPL1-like kinase, breast-tumor-amplified kinase, serine/threonine kinase 15, serine/threonine kinase 6, serine/threonine protein kinase 15, serine/threonine protein kinase 6, OTTHUMP00000031340, OTTHUMP00000031342, OTTHUMP00000031343, OTTHUMP00000031344, OTTHUMP00000031345, OTTHUMP00000166071

Official Symbol: AURKA

Accession Number(s): NP_003591.2

Human GeneID(s): [6790](#)

Important Comments: Reported variants represent identical protein: NP_003591.2; NP_940835.1; NP_940836.1; NP_940837.1; NP_940838.1; NP_940839.1

Immunogen

Peptide with sequence C-QNKESASKQS, from the C Terminus of the protein sequence according to NP_003591.2.

Please note the [peptide](#) is available for sale.

Purification and Storage

Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity chromatography using the immunizing peptide.

Supplied at 0.5 mg/ml in Tris saline, 0.02% sodium azide, pH7.3 with 0.5% bovine serum albumin.

Aliquot and store at -20°C. Minimize freezing and thawing.

Applications Tested

Peptide ELISA: antibody detection limit dilution 1:128000.

Western blot: Approx 45kDa band observed in lysates of cell line HeLa and Jurkat lysate (calculated MW of 45.8kDa according to NP_003591.2). Recommended concentration: 0.1-0.3µg/ml.

Immunoprecipitation: This antibody was deemed fit for IP under native conditions (observations from a customer).

Additional validation: This antibody has been successfully used in the following paper: Sikorski et al. (2018) PMID: 30377371.

Species Reactivity

Tested: Human

Expected from sequence similarity: Human

Specific Reference

This antibody has been successfully used in the following paper:

Krzysztof Sikorski, Adi Mehta, Marit Inngjerdengen, Flourina Thakor, Simon Kling, Tomas Kalina, Tuula A. Nyman, Maria Ekman Stensland, Wei Zhou, Gustavo A. De Souza, Lars Holden, Jan Stuchly, Markus Templin and Fridtjof Lund-Johansen

A high-throughput pipeline for validation of antibodies

Nat Methods. 2018 Nov;15(11):909-912

PMID: 30377371



EB05035 (0.1 µg/ml) staining of Jurkat lysate (35 µg protein in RIPA buffer). Primary incubation was 1 hour.
Detected by chemiluminescence.