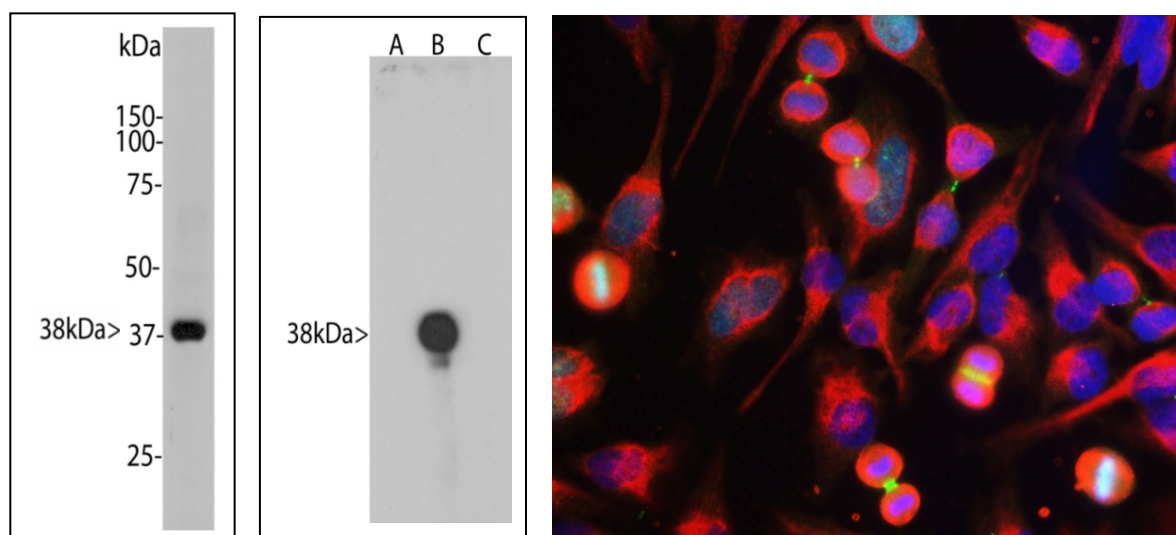


Catalogue# MCA-3F11: Mouse Monoclonal Antibody to Aurora B Kinase

The Immunogen: Aurora proteins are a family of serine/threonine protein kinases that play a key role in the regulation of cell division. The first Aurora kinase was discovered in *Drosophila* (1). Mutations of this kinase cause monopolar spindles surrounded by kinase, and the appearance of this was reminiscent of the Aurora borealis at the poles of the earth (1). Mammalian genomes encode 3 Aurora kinases named Aurora A, Aurora B, and Aurora C. All 3 contain a regulatory domain at the N terminus which is quite different between the molecules followed by a catalytic serine/threonine kinase domain which is almost identical between them. To download a sequence alignment of the 3 human Aurora proteins go to <http://encorbio.com/Alignments/Aurora%20alignment.pdf>. As a consequence antibodies raised against one Aurora family member frequently cross-react with other family members. There is a short C-terminal peptide which is also variable between the three molecules (2). Aurora A is required for centrosome duplication, entry into mitosis, formation of bipolar spindle and mitotic checkpoint (3). Aurora B is a chromosomal passenger protein and essential for chromosome condensation, kinetochore functions, spindle checkpoint activation and cytokinesis completion (4). Aurora C is heavily expressed in testis and is involved in spermatogenesis, but is also expressed in many cell lines and cancer cells and has been less well studied to date (5). Aurora A is first associated with centrosomes and then with spindle microtubules whereas Aurora B localizes to the spinal midzone and finally accumulates at the midbody. MCA-3F11 was raised against full length recombinant human Aurora B expressed in and purified from *E. coli*. The antibody was tested for binding to expressed human Aurora A, B and C and shown to react with aurora B specifically (Blot image). The HGNC name for Aurora B is AURKB.



Left: Western blot analysis of MCA-3F11 in HeLa cells. Blot of HeLa cells treated with 100ng/ml nocodazole for 18 hours was probed with MCA-3F11. Nocodazole is a microtubule polymerization inhibitor which induces cells to halt at the G2/M phase and also induces Aurora B expression. The MCA-3F11 monoclonal binds strongly to aurora B at 38 kDa. **Middle:** Blot of recombinant human Aurora A, B and C were probed with MCA-3F11. This antibody binds specifically to Aurora B. This is also consistent with the immunocytochemical staining of midzones and midbodies on HeLa cells. **Right:** HeLa cell cultures were stained with MCA-3F11 antibody (green). Aurora B stains midzones in anaphase and midbodies between the two daughter cells during telophase. It is therefore a useful marker of dividing cells. Cells were counterstained with our chicken polyclonal antibody to Vimentin **CPCA-Vim** in red. Blue is a DNA stain.

Antibody characteristics: MCA-3F11 is a mouse IgG2a class antibody with a κ light chain. MCA-3F11 recognizes aurora B kinases in western blots and in immunocytochemical experiments.

Suggestions for use: The antibody is protein G purified from tissue culture supernatant and is diluted in phosphate buffered saline at 1 mg/mL. The preparation contains 10 mM sodium azide preservative (Link to <http://www.encorbio.com/MSDS/azide.htm> for a material safety data sheet (MSDS)). The antibody solution can be used at dilutions 1:500-1:1,000 for immunofluorescence. For western blots try at 1:1,000.

Storage Instructions: Shipped on ice. Please store at 4°C for regular uses. For long term storage, please leave frozen at -20°C and avoid freeze/thaw cycles.

Limitations: This product is for research use only and is not approved for use in humans or in clinical diagnosis

References:

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4. Andrew PD, Knatko E, Moore WJ, Swedlow JR. Mitotic mechanics: the auroras come into view. *Curr Opin Cell Biol.*15(6):672-83 (2003). Review.
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