

Monoclonal anti-human ALDOA antibody (clone AT3F9)

Mouse IgG_{2a}, κ

Cat. No. IBATGA0341

Immunogen: Recombinant human ALDOA (1-364aa) purified from *E. coli*

NCBI Accession No.: NP_908930

Isotype: Mouse IgG_{2a} heavy chain and κ light chain

Clone: Anti-human ALDOA(Aldolase A) mAb, clone AT3F9, is derived from hybridization of mouse F0 myeloma cells with spleen cells from BALB/c mice immunized with a recombinant human ALDOA protein.

Description: Aldolase (fructose bisphosphate aldolase), a glycolytic enzyme, catalyzes the conversion of fructose 1, 6-bisphosphate to 3-phosphoglyceraldehyde. This ubiquitous enzyme is present as three different isozymes: aldolase A, aldolase B, and aldolase C. Research studies suggest that aldolase A expression potentially differentiates between nonneoplastic liver diseases and hepatocarcinoma. Aldolase A is found in the developing embryo and is produced in even greater amounts in adult muscle. Aldolase A expression is repressed in adult liver, kidney, and intestine and similar to aldolase C levels in brain and other nervous tissue.

Concentration: 1 mg/ml

Form: Liquid. In Phosphate-Buffered Saline (pH 7.4) with 0.02% Sodium Azide, 10% Glycerol

Storage: Can be stored at +4C. For long term storage, aliquot and store at -20C. Avoid repeated freezing and thawing cycles.

Usage: The antibody has been tested by ELISA, Western blot analysis, Flow cytometry and ICC/IF to assure specificity and reactivity. Since application varies, however, each investigation should be titrated by the reagent to obtain optimal results. Recommended dilution range for Western blot analysis is 1:1000. Recommended dilution range for ICC/IF and Flow cytometry is 1:200.

Application: ELISA, WB, ICC/IF, Flow cyt

For research use only. This product is not intended or approved for human, diagnostics or veterinary use.



Manufactured for:
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Product information

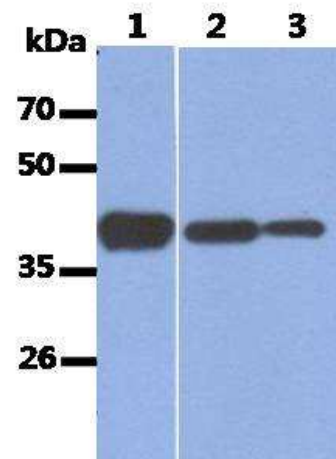
Western blot analysis

The Recombinant Human ALDOA and Cell lysates (40ug) were resolved by SDS-PAGE, transferred to PVDF membrane and probed with anti-human ALDOA antibody (1:1000). Proteins were visualized using a goat anti-mouse secondary antibody conjugated to HRP and an ECL detection system.

Lane 1.: Recombinant Human ALDOA

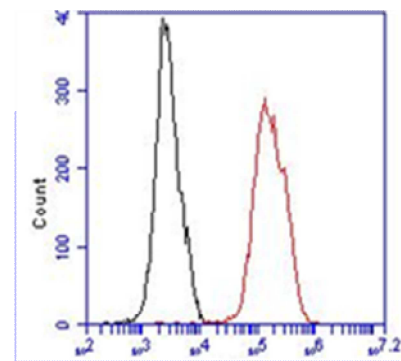
Lane 2.: HeLa cell lysate

Lane 3.: HepG2 cell lysate

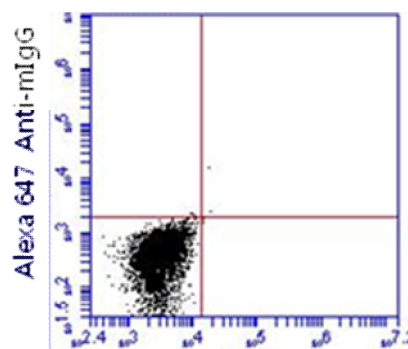


Flow cytometry

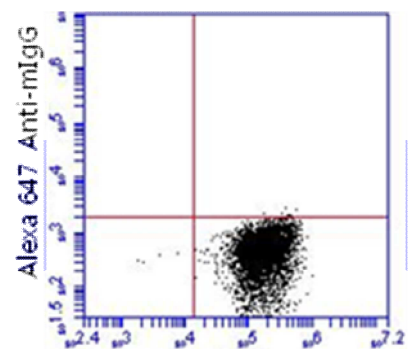
Flow cytometry analysis of ALDOA in A549 cell line, staining at 2-5ug for 1×10^6 cells (red line). The secondary antibody used goat anti-mouse IgG Alexa fluor 488 conjugate. Isotype control antibody was mouse IgG (black line).



Alexa 488 Anti-Aldolase A



Alexa 488 Anti-mIgG



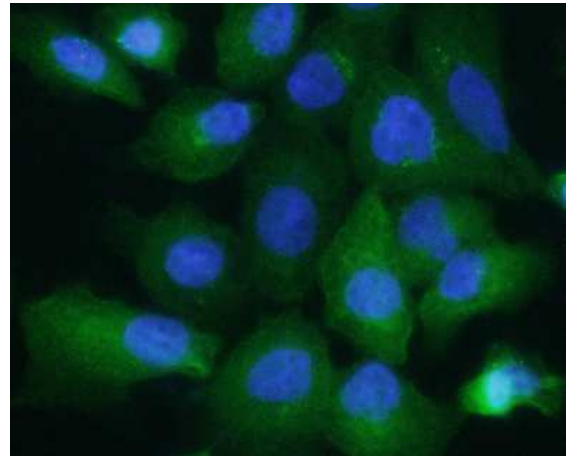
Alexa 488 Anti-Aldolase A

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ICC/IF analysis

ICC/IF analysis of ALDOA in A549 cells, stained with DAPI (Blue) for nucleus staining and monoclonal anti-human A549 antibody (1:200) with goat anti-mouse IgG-Alexa fluor 488 conjugate (Green).



General references: Castaldo G., *et al.*(2000) *Clin Chem.* **46(7)**: 901-906
Long F., *et al.*(2014) *Oncol Rep.* **32(5)**: 2031-2037
Caspi M., *et al.*(2014) *Mol Cancer.* **13**: 164

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