

## Monoclonal anti-human 14-3-3 tau antibody (clone AT1A1)

Mouse IgG<sub>1</sub>, κ

Cat. No. IBATGA0132

**Immunogen:** Recombinant human 14-3-3 tau (1-245aa) purified from *E. coli*

**NCBI Accession No.:** NP\_006817

**Isotype:** Mouse IgG<sub>1</sub> heavy chain and κ light chain

**Clone:** Anti-human 14-3-3 tau mAb, clone AT1A1, is derived from hybridization of mouse F0 myeloma cells with spleen cells from BALB/c mice immunized with a recombinant human 14-3-3 tau protein.

**Description:** The 14-3-3 family of proteins plays a key regulatory role in signal transduction, checkpoint control, apoptotic and nutrient-sensing pathways. 14-3-3 proteins are highly conserved and ubiquitously expressed. There are at least seven isoforms, β, γ, ε, σ, ζ, τ and η that have been identified in mammals. The 14-3-3 tau, a subtype of the 14-3-3 family of proteins, was found in T Cells, brain and testes. This 14-3-3 tau is upregulated in patients with amyotrophic lateral sclerosis.

**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, ICC/IF and Flow cytometry to assure specificity and reactivity. Since application varies, however, each investigation should be titrated by the reagent to obtain optimal results.

**Application:** ELISA, WB, ICC/IF, Flow cytometry

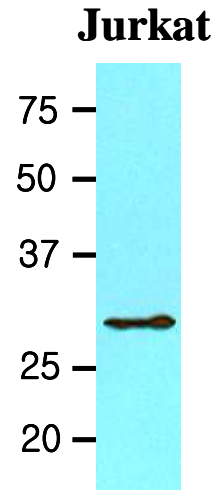
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# Product information



## Western blot analysis

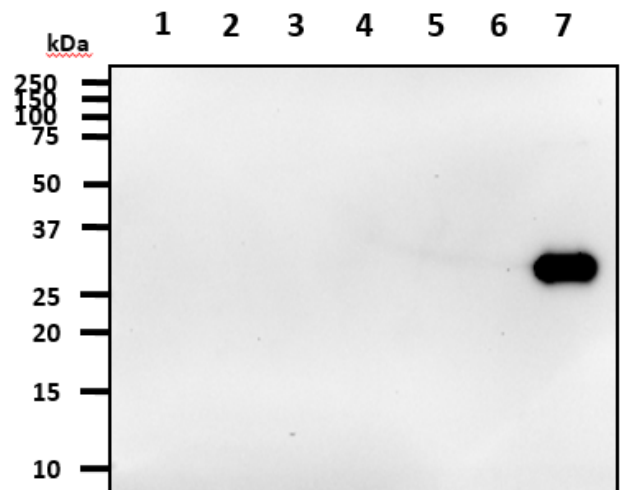
Cell lysates of Jurkat (20ug) was resolved by SDS-PAGE, transferred to NC membrane and probed with anti-human 14-3-3 tau antibody (1:1000). Proteins were visualized using a goat anti-mouse secondary antibody conjugated to HRP and an ECL detection system.



## Western blot analysis

The recombinant proteins (50ng) were resolved by SDS-PAGE, transferred to PDVF membrane and probed with anti-human 14-3-3 tau antibody (1:1000). Proteins were visualized using a goat anti-mouse secondary antibody conjugated to HRP and an ECL detection system.

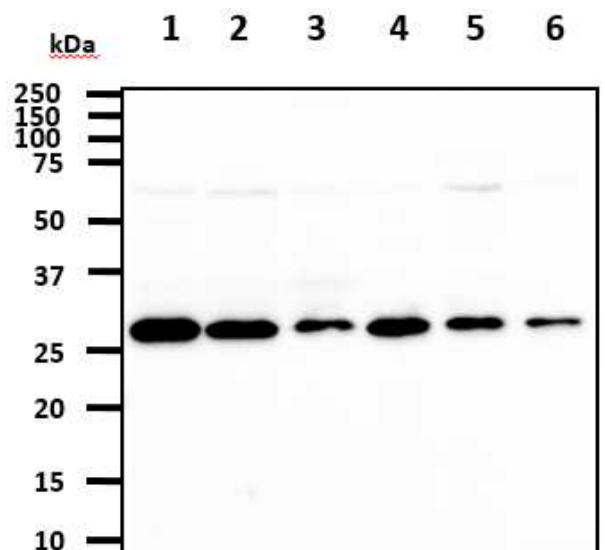
- Lane 1. : Recombinant Human YWHAZ
- Lane 2. : Recombinant Human YWHAB
- Lane 3. : Recombinant Human YWHAE
- Lane 4. : Recombinant Human YWHAH
- Lane 5. : Recombinant Human YWHAG
- Lane 6. : Recombinant Human SFN
- Lane 7. : Recombinant Human YWHAQ



## Western blot analysis

The cell lysates (40ug) were resolved by SDS-PAGE, transferred to PDVF membrane and probed with anti-human 14-3-3 tau antibody (1:1000). Proteins were visualized using a goat anti-mouse secondary antibody conjugated to HRP and an ECL detection system.

- Lane 1. : MDA-MB-21 cell lysate
- Lane 2. : HeLa cell lysate
- Lane 3. : A431 cell lysate
- Lane 4. : NIH3T3 cell lysate
- Lane 5. : 293T cell lysate
- Lane 6. : HepG2 cell lysate



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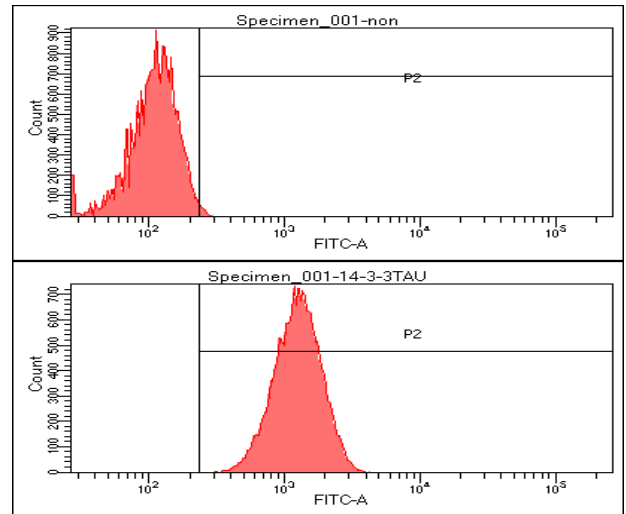


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# Product information

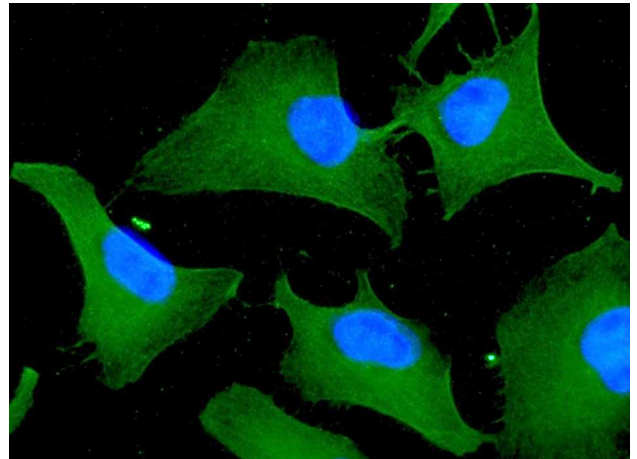
## Flow cytometry

Flow cytometry analysis of 14-3-3 tau in A549 cell line, staining at 2-5ug for  $1 \times 10^6$  cells. The secondary antibody used goat anti-mouse IgG Alexa fluor 488 conjugate.



## ICC/IF analysis

ICC/IF analysis of 14-3-3 tau in HeLa cell line, stained with DAPI (Blue) for nucleus staining and monoclonal anti-human 14-3-3 tau antibody (1:100) with goat anti-mouse IgG-Alexa fluor 488 conjugate (Green).



**General references:** Liu YC., *et al.* (1996) *J Biol Chem.* **271**: 14591-14595.  
Xiao B., *et al.* (1995) *Nature.* **376**: 188-191.

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