

## Monoclonal anti-human FADD antibody (clone J1D2)

Mouse IgG<sub>2b</sub>, κ

Cat. No. IBAFA0901

**Immunogen:** Recombinant human FADD (1-208aa) purified from *E. coli*.

**NCBI Accession No.:** NP\_003815

**Isotype:** Mouse IgG<sub>2b</sub> heavy chain and κ light chain

**Clone:** Anti-human FADD mAb, clone J1D2, is derived from hybridization of mouse FO myeloma cells with spleen cells from BALB/c mice immunized with a recombinant human FADD protein.

**Description:** FADD (Fas-associated protein with death domain) is an adaptor molecule that interacts with various cell surface receptors and mediates cell apoptotic signals. This protein is implicated in survival/proliferation and cell cycle progression. FADD functions are also regulated via cellular sublocalization, protein phosphorylation, and inhibitory molecules.

**Concentration:** 1mg/ml

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

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

**Usage:** The antibody has been tested by ELISA, Western blot analysis, ICC/IF, Flow cytometry and immunohistochemistry analysis 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 cyt, IHC

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



Manufactured for:

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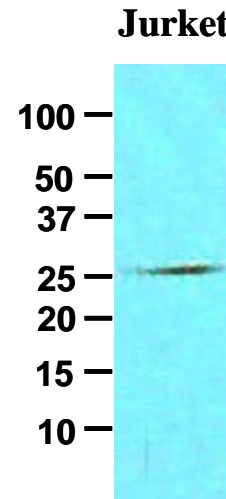
Email: [info@ibl-america.com](mailto:info@ibl-america.com)

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# Product information

## Western blot analysis

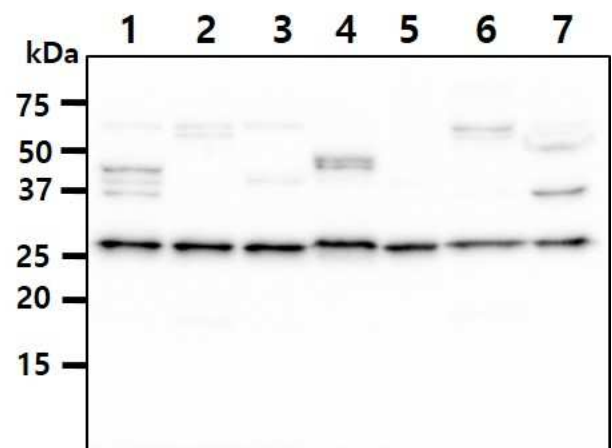
Cell lysates of Jurkat (30ug) were resolved by SDS-PAGE, transferred to NC membrane and probed with anti-human FADD (1:500). Proteins were visualized using a goat anti-mouse secondary antibody conjugated to HRP and an ECL detection system.



## Western blot analysis

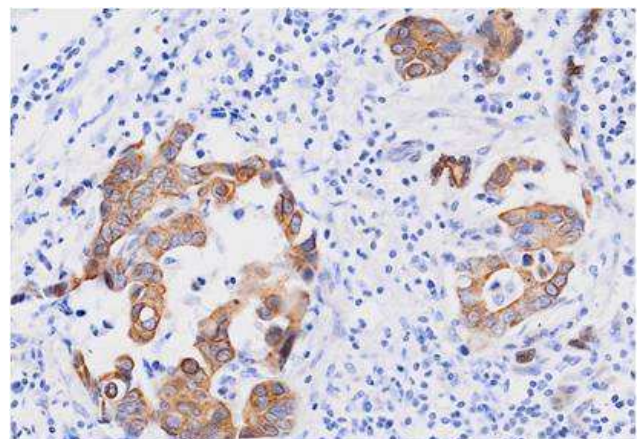
The Cell lysates (40ug) were resolved by SDS-PAGE, transferred to PVDF membrane and probed with anti-human FADD antibody (1:500). Proteins were visualized using a goat anti-mouse secondary antibody conjugated to HRP and an ECL detection system.

- Lane 1. : HeLa cell lysate
- Lane 2. : Raw264.7 cell lysate
- Lane 3. : MCF7 cell lysate
- Lane 4. : A431 cell lysate
- Lane 5. : Ramos cell lysate
- Lane 6. : Raji cell lysate
- Lane 7. : Balb/3T3 cell lysate



## Immunohistochemistry

Paraffin embedded sections of human breast cancer tissue were incubated with anti-human FADD (1:50) for 2 hours at room temperature. Antigen retrieval was performed in 0.1M sodium citrate buffer and detected using Diaminobenzidine (DAB)

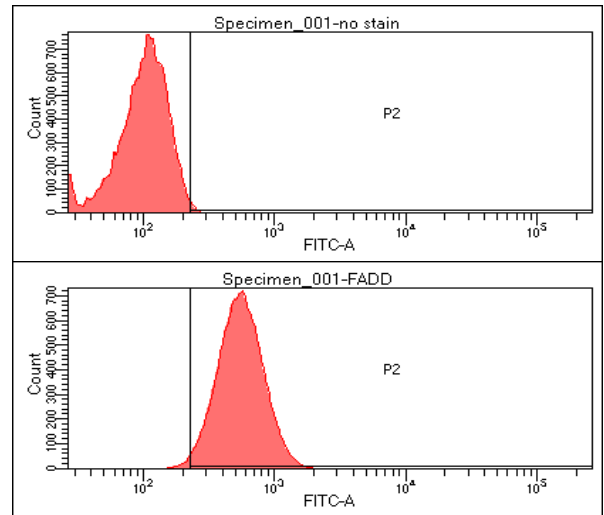


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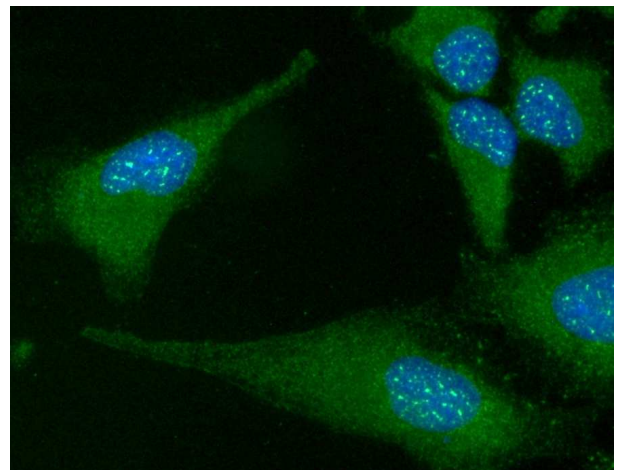
## Flow cytometry

Flow cytometry analysis of FADD in HeLa 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 FADD in HeLa cells line, stained with DAPI (Blue) for nucleus staining and monoclonal anti-human FADD antibody (1:100) with goat anti-mouse IgG-Alexa fluor 488 conjugate (Green).



- General references:**
- Léa Tourneur., *et al.* (2005) *Medical Immunology*. **4**:1.
  - Tsao, C.H., *et al.* (2008) *J. Gen. Virol.* **89**; (PT 8), 1930-1941.
  - Douglas D. Bannerman, *et al.* (2002). *J. Clin. Invest.* **109**:419–425.

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