Anti-Caspase-8 Antibody [6G1]

RT1099



Product Type: Mouse monoclonal IgG1, primary antibodies

Species reactivity: Human

Applications: WB, IP, IF, IHC-P

Molecular Wt: 55kDa
Clone number: 6G1

Description: Initiator caspases, which include caspase-8, activate effector caspases by cleaving inactive

forms of effector caspases. In the activation cascade responsible for apoptosis induced by TNFRSF1A and mediated by TNFRSF6/FAS, caspase-8 is the most upstream protease. Caspase-8 binds to adaptor molecule FADD, forming an aggregate referred to as death-inducing signaling complex (DISC), which activates caspase-8. The actived protein is released from the complex and further activates downstream apoptotic proteases. Caspase-8, which is a heterodimer consisting of two subunits (p18 and p10), is widely expressed, but is detected at highest levels in peripheral blood leukocytes (PBLs), thymus, liver and spleen. Defects in CASP8, the gene encoding for caspase-8, may cause CASP8D (caspase-8 deficiency disorder), which is characterized by splenomegaly and CD95-induced apoptosis of PBLs, may lead to immunodeficiency due to defects in T lymphocyte, NK cell

and B lymphocyte activation.

Immunogen: peptide

Positive control: Hela, Jurkat, CCRF-CEM, MOLT-4

Subcellular location: Cytoplasm

Database links: SwissProt: Q14790 Human

Recommended Dilutions:

WB 1:100-1:1,000

IP 1-2 μg per 100-500 μg of total protein(1 ml of cell lysate)

IF 1:50-500 IHC-P 1:50-500

Storage Buffer: 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Storage Instruction: Store at +4 ℃

Purity: Protein A affinity purified.

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Images

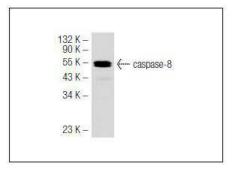


Fig1: Western blot analysis of caspase-8 expression in Jurkat whole cell lysate.

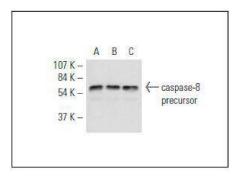


Fig2: Western blot analysis of caspase-8 expression in CCRF-CEM (A), HeLa (B) and MOLT-4 (C) whole cell lysates.

Note: All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE".

Background References

- 1. Wang, O., et al. 2011. Anticancer activity of 2α, 3α, 19β, 23β-Tetrahydroxyurs-12-en-28-oic acid (THA), a novel triterpenoid isolated from Sinojackia sarcocarpa. PLoS ONE 6: e21130.
- 2. Cleveland, J.L., et al. 1995. Contenders in FAS-L/TNF death signaling. Cell 81: 479-482.