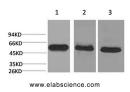
# Elabscience®

## **CASP8 Monoclonal Antibody**

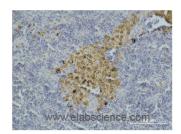
Catalog No.E-AB-22107ReactivityH,M,RStorageStore at -20°C. Avoid freeze / thaw cycles.HostMouseApplicationsWB,IHC-p,IFIsotypeIgG

Note: Centrifuge before opening to ensure complete recovery of vial contents.

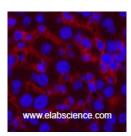
### **Images**



Western Blot analysis of 1) Hela, 2) Mouse brain, 3) Rat brain with CASP8 Monoclonal Antibody



Immunohistochemistry of paraffinembedded Mouse spleen tissue with CASP8 Monoclonal Antibody



Immunofluorescence analysis of Mouse liver tissue using CASP8 Monoclonal Antibody at dilution of 1:200.

### **Immunogen Information**

Immunogen Recombinant Protein

Swissprot Q14790

**Synonyms** CASP8,MCH5,Caspase-8,CASP-8,Apoptotic cysteine

protease, Apoptotic protease Mch-5, CAP4, FADD-homologous ICE/ced-3-like protease, FADD-like

ICE,FLICE,ICE-like apoptotic protease 5,MORT1-associated ced-3 homolog,MACH

#### **Product Information**

Observed MW 43,57kDa

**Buffer** PBS with 0.02% sodium azide and 50% glycerol pH

7.4.

**Purify** Protein A purification

Clone No. Clone:2E3

**Dilution** WB 1:500-1:2000, IHC 1:100-1:300, IF 1:100-1:300

#### **Background**

Most upstream protease of the activation cascade of caspases responsible for the TNFRSF6/FAS mediated and TNFRSF1A induced cell death. Binding to the adapter molecule FADD recruits it to either receptor. The resulting aggregate called death-inducing signaling complex (DISC) performs CASP8 proteolytic activation. The active dimeric enzyme is then liberated from the DISC and free to activate downstream apoptotic proteases. Proteolytic fragments of the N-terminal propeptide (termed CAP3, CAP5 and CAP6) are likely retained in the DISC. Cleaves and activates CASP3, CASP4, CASP6, CASP7, CASP9 and CASP10. May participate in the GZMB apoptotic pathways. Cleaves ADPRT. Hydrolyzes the small-molecule substrate, Ac-Asp-Glu-Val-Asp-AMC. Likely target for the cowpox virus CRMA death inhibitory protein. Isoform 5, isoform 6, isoform 7 and isoform 8 lack the catalytic site and may interfere with the pro-apoptotic activity of the complex.

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