

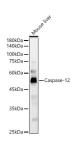
Tel:240-252-7368(USA) Fax: 240-252-7376(USA) techsupport@elabscience.com Website: www.elabscience.com

# **Caspase-12 Polyclonal Antibody**

Catalog No.E-AB-64000ReactivityH,M,RStorageStore at -20°C. Avoid freeze / thaw cycles.HostRabbitApplicationsWB,IHCIsotypeIgG

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

## **Images**



Western blot analysis of Mouse liver using Caspase-12 Polyclonal Antibody at 1:500 dilution.



Immunohistochemistry of paraffinembedded human tonsil using Caspase-12 Polyclonal Antibody at dilution of 1:100 (40x lens). Perform high pressure antigen retrieval with 10 mM citrate buffer pH 6.0 before commencing with IHC staining protocol.



Immunohistochemistry of paraffinembedded mouse spleen using Caspase-12 Polyclonal antibody at dilution of 1:100 (40x lens).Perform high pressure antigen retrieval with 10

# **Immunogen Information**

**Immunogen** A synthetic peptide of human Caspase-12

 GeneID
 100506742

 Swissprot
 Q6UXS9

Synonyms CASP12,CASP-12,CASP12P1

#### **Product Information**

Calculated MW 19kDa/29kDa/38kDa

Observed MW 55KDa

**Buffer** PBS with 0.05% proclin300,50% glycerol,pH7.3.

**Purify** Affinity purification

**Dilution** WB 1:500-1:2000,IHC 1:50-1:200

### **Background**

Caspases are cysteine proteases that cleave C-terminal aspartic acid residues on their substrate molecules. This gene is most highly related to members of the ICE subfamily of caspases that process inflammatory cytokines. In rodents, the homolog of this gene mediates apoptosis in response to endoplasmic reticulum stress. However, in humans this gene contains a polymorphism for the presence or absence of a premature stop codon. The majority of human individuals have the premature stop codon and produce a truncated non-functional protein. The read-through codon occurs primarily in individuals of African descent and carriers have endotoxin hypo-responsiveness and an increased susceptibility to severe sepsis. Several alternatively spliced transcript variants have been noted for this gene. [provided by RefSeq, Feb 2011]

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