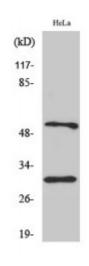


Anti-Caspase-10 B/C antibody



Description

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Rabbit polyclonal to Caspase-10 B/C.

Model STJ92018

Host Rabbit

Reactivity Human

Applications ELISA, IF, IHC, WB

Immunogen Synthesized peptide derived from human Caspase-10 B/C

Immunogen Region 400-480 aa, C-terminal

Gene Symbol <u>CASP10</u>

Dilution range WB 1:500-1:2000IHC 1:100-1:300IF 1:200-1:1000ELISA 1:10000

Specificity Caspase-10 B/C Polyclonal Antibody detects endogenous levels of

Caspase-10 B/C protein.

Tissue Specificity Detectable in most tissues. Lowest expression is seen in brain, kidney,

prostate, testis and colon.

Purification The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Note For Research Use Only (RUO).

Protein Name Caspase-10 CASP-10 Apoptotic protease Mch-4 FAS-associated death

domain protein interleukin-1B-converting enzyme 2 FLICE2 ICE-like apoptotic protease 4 Caspase-10 subunit p23/17 Caspase-10 subunit p12

Molecular Weight 60 kDa

Clonality Polyclonal

Conjugation Unconjugated

Isotype IgG

Formulation Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Concentration 1 mg/ml

Storage Instruction Store at -20°C, and avoid repeat freeze-thaw cycles.

Alternative Names Caspase-10 CASP-10 Apoptotic protease Mch-4 FAS-associated death

domain protein interleukin-1B-converting enzyme 2 FLICE2 ICE-like apoptotic protease 4 Caspase-10 subunit p23/17 Caspase-10 subunit p12

Function Involved in the activation cascade of caspases responsible for apoptosis

execution. Recruited to both Fas- and TNFR-1 receptors in a FADD dependent manner. May participate in the granzyme B apoptotic pathways. Cleaves and activates caspase-3, -4, -6, -7, -8, and -9. Hydrolyzes the small-molecule substrates, Tyr-Val-Ala-Asp-|-AMC and Asp-Glu-Val-Asp-|-AMC.

Isoform 7 can enhance NF-kappaB activity but promotes only slight apoptosis.

Isoform C is proteolytically inactive.

Post-translational Cleavage by granzyme B and autocatalytic activity generate the two active

Modifications subunits.

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