Recombinant Protein Technical Manual

Recombinant Human cIAP1/HIAP2 Protein (AVI Tag)(Active)

RPES3687

Product Data:

Product SKU: RPES3687

Size: 50µg

Species: Human

Expression host: E. coli

Uniprot: NP_001157.1

Protein Information:

Molecular Mass: 26.5 kDa

AP Molecular Mass: 26.5 kDa

Tag:

Bio-activity: Measured by its ability to inhibit DEVD-AFC cleavage activity in cell extracts activated by addition of cytochrome c and dATP. The IC50 for this effect is typically 25-750 nM.

Purity: > 92 % as determined by reducing SDS-PAGE.

Endotoxin: Please contact us for more information.

Storage: Lyophilized proteins are stable for up to 12 months when stored at -20 to -80°C. Reconstituted protein solution can be stored at 4-8℃ for 2-7 days. Aliquots of reconstituted samples are stable at < -20℃ for 3 months.

Shipping: This product is provided as lyophilized powder which is shipped with ice packs.

Formulation: Lyophilized from sterile 10mM Tris, 5% glycerol, 0.5mM EDTA, 5mM DTT, pH 7.5

Reconstitution: Please refer to the printed manual for detailed information.

Application:

Synonyms: API1;c-IAP1;cIAP1;Hiap-2;HIAP2;MIHB;RNF48

Immunogen Information:

Sequence: Glu 144-Leu 356

Background:

The cellular inhibitor of apoptosis protein (cIAP1) is a member of the Inhibitor of Apoptosis family proteinsare (IAP) whose members are characterized by a novel domain of about 70 amino acids termed baculoviral IAP repeats (BIRs). The BIR domains of cIAP1 and cIAP2 bind to caspases, the key effector proteases of apoptosis. The IAP protein family which can enhance cell survival are crucial regulators of programmed cell death. Both cIAP1 and cIAP2 are the E3 ubiquitin protein isopeptide ligases for Smac, taking part in promoting cancer survival through functioning as E3 ubiqitin ligases. Removal of cIAP1 by genetic deletion may result in NF-κB signaling activation that induces TNFα production and in killing sensitive tumor cells through enhanced TNF-R1 death-receptor signaling and caspase 8 activation. The substrate-dependent E3 activity of cIAPs is mediated by their RING domains and is dependent on the specific interactions between cIAPs and Smac. cIAP1 and cIAP2 are also reported to be regulators of NF-kB activation upon TNFαtreatment.