

# Recombinant Protein Technical Manual

# Recombinant Human XIAP/BIRC4 Protein (AVI Tag)(Active) RPES3648

#### Product Data:

**Product SKU:** RPES3648 **Size:** 50μg

Species: Human Expression host: E. coli

**Uniprot:** NP 001158.2

#### **Protein Information:**

Molecular Mass: 29.1 kDa

AP Molecular Mass: 29.1 kDa

Tag:

**Bio-activity:** Measured by its binding ability in a functional ELISA. Immobilized recombinant

human SMAC-His at 10 μg/ml (100 μl/well) can bind recombinant human XIAP-AVI

with a linear range of 0.125.0 μg/ml.

**Purity:** > 75 % 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°C for 2-7 days. Aliquots of

reconstituted samples are stable at < -20°C for 3 months.

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

**Formulation:** Lyophilized from sterile 25mM Tris, 10mM DTT, 1% glycerol, 0.2M Glutamine

Potassium, pH 8.0

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

**Application:** Functional ELISA

**Synonyms:** API3;BIRC4;hIAP-3;hIAP3;IAP-3;ILP1;MIHA;XLP2

## **Immunogen Information:**

Sequence: Leu 121-Thr 356

## Background:

E3 ubiquitin-protein ligase XIAP / BIRC4, also known as inhibitor of apoptosis protein 3, X-linked inhibitor of apoptosis protein, and IAP-like protein, is a protein that belongs to a family of apoptotic suppressor proteins. Members of this family share a conserved motif termed, baculovirus IAP repeat, which is necessary for their anti-apoptotic function. XIAP / BIRC4 functions through binding to tumor necrosis factor receptor-associated factors TRAF1 and TRAF2 and inhibits apoptosis induced by menadione, a potent inducer of free radicals, and interleukin 1-beta converting enzyme. XIAP / BIRC4 also inhibits at least two members of the caspase family of cell-death proteases, caspase-3 and caspase-7. Mutations in this encoding gene are the cause of X-linked lymphoproliferative syndrome. Alternate splicing results in multiple transcript variants. Thought to be the most potent apoptosis suppressor, XIAP / BIRC4, directly binds and inhibits caspases -3, -7 and -9. Survivin, which also binds to several caspases, is up-regulated in a many tumour cell types. Defects in XIAP / BIRC4 are the cause of lymphoproliferative syndrome X-linked type 2 (XLP2). XLP is a rare immunodeficiency characterized by extreme susceptibility to infection with Epstein-Barr virus (EBV). Symptoms include severe or fatal mononucleosis, acquired hypogammaglobulinemia, pancytopenia and malignant lymphoma.