

Recombinant Protein Technical Manual Recombinant Mouse FAS/TNFRSF6 Protein (His Tag)(Active) RPES0092

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Product SKU: RPES0092	Size: 50μg
Species: Mouse	Expression host: HEK293 Cells

Uniprot: NP_032013.2

Protein Information

Molecular Mass:	18 kDa
AP Molecular Mass:	
Tag:	C-His
Bio-activity:	Measured by its ability to inhibit Fas Ligand induced apoptosis of Jurkat human acute T cell leukaemia cells. The ED50 for this effect is typically 1-8 μg/mL in the presence of 20 ng/mL recombinant human Fas ligand.
Purity:	> 98 % as determined by SDS-PAGE
Endotoxin:	< 1.0 EU per μg of the protein as determined by the LAL method.
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 PBS, pH 7.4
Reconstitution:	Please refer to the printed manual for detailed information.
Application:	
Synonyms:	AI196731 Protein, APO1 Protein, APT1 Protein, CD95 Protein, lpr Protein, TNFR6 Protein, Tnfrsf6 Protein

Sequence: Met 1-Arg 169

Background:

CD95 (APO/Fas) is an important inducer of the extrinsic apoptosis signalling pathway and therapy induced apoptosis of many tumor cells has been linked to the activity of CD95. is a prototype death receptor characterized by the presence of an 80 amino acid death domain in its cytoplasmic tail. This domain is essential for the recruitment of a number of signalling components upon activation by either agonistic anti-CD95 antibodies or cognate CD95 ligand that initiate apoptosis. The complex of proteins that forms upon triggering of CD95 is called the death-inducting signalling complex (DISC). The DISC consists of an adaptor protein and initiator caspases and is essential for induction of apoptosis. CD95 is also crucial for the negative selection of B cells within the germinal center (GC). Impairment of CD95-mediated apoptosis results in defective affinity maturation and the persistence of autoreactive B-cell clones. Changes in the expression of CD95 and/or its ligand CD95L are frequently found in human cancer. The downregulation or mutation of CD95 has been proposed as a mechanism by which cancer cells avoid destruction by the immune system through reduced apoptosis sensitivity. Thus, CD95 has therefore been viewed as a tumor suppressor. CD95 has been reported to be involved in the activation of NF-kappaB, MAPK3/ERK1, MAPK8/JNK, and the alternate pathways for CTL-mediated cytotoxicity. Accordingly, this protein is implicated in the pathogenesis of various malignancies and diseases of the immune system. The CD95/CD95L system was implicated in the etiology of inflammatory bowel disease (IBD) based, primarily, on the finding that CD95 is highly expressed in the intestinal epithelial cells and that epithelial apoptosis is increased in IBD.