# abcam

### Product datasheet

## Recombinant human Fas protein ab 199806

**Description** 

Product name Recombinant human Fas protein

**Biological activity** Fully biologically active when compared to standard. The ED<sub>50</sub> as determined by its ability to

inhibit the cytotoxicity of Jurkat cells is between 10-15  $\mu$ g/mL in the presence of 2  $\eta$ g/mL of

rHuFas Ligand.

Purity > 95 % SDS-PAGE.

assessed also by HPLC analysis

**Endotoxin level** < 1.000 Eu/µg

**Expression system** Escherichia coli

Accession P25445

Protein length Protein fragment

Animal free No

Nature Recombinant

**Species** Human

Sequence RLSSKSVNAQ VTDINSKGLE LRKTVTTVET

QNLEGLHHDG QFCHKPCPPG ERKARDCTVN
GDEPDCVPCQ EGKEYTDKAH FSSKCRRCRL
CDEGHGLEVE INCTRTQNTK CRCKPNFFCN

STVCEHCDPC TKCEHGIIKE CTLTSNTKCK EEGSRSN

Predicted molecular weight 18 kDa

Amino acids 17 to 173

**Specifications** 

Our **Abpromise guarantee** covers the use of **ab199806** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications HPLC

SDS-PAGE

**Functional Studies** 

Form Lyophilized

**Preparation and Storage** 

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Stability and Storage Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle. Reconstitute for long term storage.

pH: 7.40

Constituent: 100% PBS

This product is an active protein and may elicit a biological response in vivo, handle with caution.

Reconstitution Lyophilized from a 0.2 µm filtered concentrated solution. We recommend that this vial be briefly

centrifuged prior to opening to bring the contents to the bottom. Reconstitute in sterile distilled water or aqueous buffer containing 0.1 % BSA to a concentration of 0.1-1.0 mg/mL. Stock solutions should be apportioned into working aliquots and stored at = -20 °C. Further dilutions

should be made in appropriate buffered solutions.

#### **General Info**

Function Receptor for TNFSF6/FASLG. The adapter molecule FADD recruits caspase-8 to the activated

receptor. The resulting death-inducing signaling complex (DISC) performs caspase-8 proteolytic activation which initiates the subsequent cascade of caspases (aspartate-specific cysteine proteases) mediating apoptosis. FAS-mediated apoptosis may have a role in the induction of peripheral tolerance, in the antigen-stimulated suicide of mature T-cells, or both. The secreted

isoforms 2 to 6 block apoptosis (in vitro).

**Tissue specificity** Isoform 1 and isoform 6 are expressed at equal levels in resting peripheral blood mononuclear

cells. After activation there is an increase in isoform 1 and decrease in the levels of isoform 6.

**Involvement in disease**Defects in FAS are the cause of autoimmune lymphoproliferative syndrome type 1A (ALPS1A)

[MIM:601859]; also known as Canale-Smith syndrome (CSS). ALPS is a childhood syndrome

involving hemolytic anemia and thrombocytopenia with massive lymphadenopathy and

splenomegaly.

**Sequence similarities**Contains 1 death domain.

Contains 3 TNFR-Cys repeats.

**Domain**Contains a death domain involved in the binding of FADD, and maybe to other cytosolic adapter

proteins.

**Cellular localization** Secreted and Cell membrane.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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