abcam

Product datasheet

Human Fas ELISA Kit ab100513

2 Images

Overview

Product name Human Fas ELISA Kit

Detection methodColorimetric

Sample type Cell culture supernatant, Serum, Plasma

Assay type Sandwich (quantitative)

Sensitivity < 5 pg/ml

Range 2.74 pg/ml - 2000 pg/ml

Recovery > 85 %

Sample specific recovery

Sample type	Average %	Range
Cell culture supernatant	91.24	82% - 105%
Serum	84.46	78% - 102%
Plasma	86.37	80% - 104%

Assay duration Multiple steps standard assay

Species reactivity Reacts with: Human

Product overview Abcam's Fas Human ELISA (Enzyme-Linked Immunosorbent Assay) kit is an in vitro enzyme-

linked immunosorbent assay for the quantitative measurement of Human Fas in serum, plasma,

and cell culture supernatants.

This assay employs an antibody specific for Human Fas coated on a 96-well plate. Standards and samples are pipetted into the wells and Fas present in a sample is bound to the wells by the immobilized antibody. The wells are washed and biotinylated anti-Human Fas antibody is added. After washing away unbound biotinylated antibody, HRP-conjugated streptavidin is pipetted to the wells. The wells are again washed, a TMB substrate solution is added to the wells and color develops in proportion to the amount of Fas bound. The Stop Solution changes the color from blue to yellow, and the intensity of the color is measured at 450 nm.

Get results in 90 minutes with Human FASLG Receptor ELISA Kit (CD95) (<u>ab183360</u>) from our SimpleStep ELISA[®] range.

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Notes Optimization may be required with urine samples

Platform Microplate

Properties

Storage instructions Store at -20°C. Please refer to protocols.

Components	1 x 96 tests
20X Wash Buffer	1 x 25ml
5X Assay Diluent B	1 x 15ml
640X HRP-Streptavidin Concentrate	1 x 200µl
Assay Diluent A	1 x 30ml
Biotinylated anti-Human Fas	2 vials
Fas Microplate (12 strips x 8 wells)	1 unit
Recombinant Human Fas Standard (lyophilized)	2 vials
Stop Solution	1 x 8ml
TMB One-Step Substrate Reagent	1 x 12ml

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.

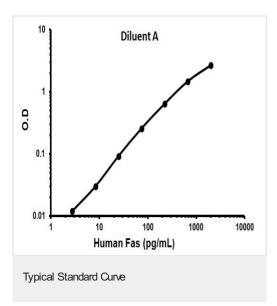
Contains 3 TNFR-Cys repeats.

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

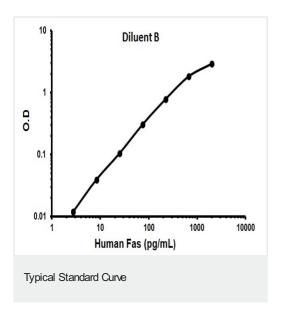
proteins.

Cellular localization Secreted and Cell membrane.

Images



Representative Standard Curve using ab100513.



Representative Standard Curve using ab100513.

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