

**Phospho-FAS(Y291) Antibody**  
**Affinity Purified Rabbit Polyclonal Antibody (Pab)**  
**Catalog # AP3310a**

**Specification**

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**Phospho-FAS(Y291) Antibody - Product Information**

Application	DB,E
Primary Accession	<a href="#">P25445</a>
Reactivity	Human
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit IgG
Calculated MW	37732

**Phospho-FAS(Y291) Antibody - Additional Information**

**Gene ID** 355

**Other Names**

Tumor necrosis factor receptor superfamily member 6, Apo-1 antigen, Apoptosis-mediating surface antigen FAS, FASLG receptor, CD95, FAS, APT1, FAS1, TNFRSF6

**Target/Specificity**

This FAS Antibody is generated from rabbits immunized with a KLH conjugated synthetic phosphopeptide corresponding to amino acid residues surrounding Y291 of human FAS.

**Dilution**

DB~~1:500

**Format**

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

**Storage**

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions**

Phospho-FAS(Y291) Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

**Phospho-FAS(Y291) Antibody - Protein Information**

**Name** FAS

**Synonyms** APT1, FAS1, TNFRSF6

**Function** Receptor for TNFSF6/FASLG. The adapter molecule FADD recruits caspase CASP8 to the

activated receptor. The resulting death-inducing signaling complex (DISC) performs CASP8 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).

#### Cellular Location

[Isoform 1]: Cell membrane; Single-pass type I membrane protein. Membrane raft [Isoform 3]: Secreted. [Isoform 5]: Secreted.

#### Tissue Location

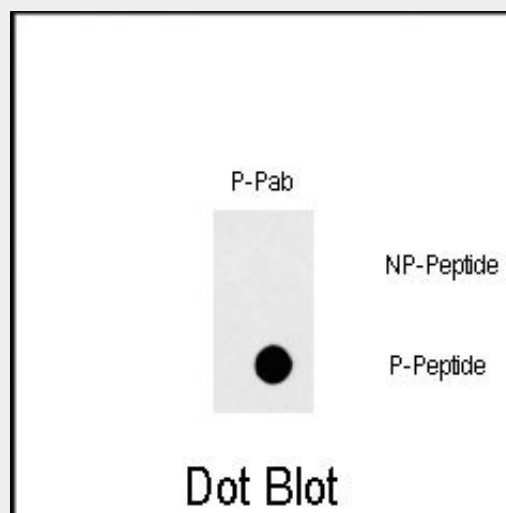
Isoform 1 and isoform 6 are expressed at equal levels in resting peripheral blood mononuclear cells. After activation there is an increase in isoform 1 and decrease in the levels of isoform 6.

### Phospho-FAS(Y291) Antibody - Protocols

Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

### Phospho-FAS(Y291) Antibody - Images



Dot blot analysis of Phospho-FAS-Y291 polyclonal antibody (Cat# AP3310a) on nitrocellulose membrane. 50ng of Phospho-peptide or Non Phospho-peptide per dot were adsorbed. Antibody working concentration was 0.5ug per ml. P-Pab: phospho-antibody; P-Peptide: phospho-peptide; NP-Peptide: non-phospho-peptide.

### Phospho-FAS(Y291) Antibody - Background

FAS is a member of the TNF-receptor superfamily. This receptor contains a death domain. It has been shown to play a central role in the physiological regulation of programmed cell death, and has

been implicated in the pathogenesis of various malignancies and diseases of the immune system. The interaction of this receptor with its ligand allows the formation of a death-inducing signaling complex that includes Fas-associated death domain protein (FADD), caspase 8, and caspase 10. The autoproteolytic processing of the caspases in the complex triggers a downstream caspase cascade, and leads to apoptosis. This receptor has been also shown to activate NF-kappaB, MAPK3/ERK1, and MAPK8/JNK, and is found to be involved in transducing the proliferating signals in normal diploid fibroblast and T cells.

#### **Phospho-FAS(Y291) Antibody - References**

- Wang, W.H., et al., Mol. Cell. Biol. 24(23):10352-10365 (2004).  
Inaba, H., et al., FEBS Lett. 43(7):729-736 (2004).  
Delmas, D., et al., Oncogene 23(55):8979-8986 (2004).  
Siegel, R.M., et al., J. Cell Biol. 167(4):735-744 (2004).  
Qiao, S., et al., FEBS Lett. 577(3):451-454 (2004).