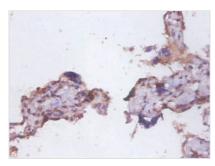






## **FAS** Antibody

CSB-PA06354A0Rb
Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
P25445
Recombinant Human Tumor necrosis factor receptor superfamily member 6 protein (26-171AA)
Rabbit
Human
ELISA, IHC, IF; Recommended dilution: IHC:1:20-1:200, IF:1:50-1:500
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).
Liquid
Non-conjugated
Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, PH 7.4
Constituents: 50% Glycerol, 0.01M PBS, PH 7.4
Constituents: 50% Glycerol, 0.01M PBS, PH 7.4 >95%, Protein G purified
Constituents: 50% Glycerol, 0.01M PBS, PH 7.4 >95%, Protein G purified  IgG
Constituents: 50% Glycerol, 0.01M PBS, PH 7.4  >95%, Protein G purified  IgG  Polyclonal  Tumor necrosis factor receptor superfamily member 6 (Apo-1 antigen) (Apoptosis-mediating surface antigen FAS) (FASLG receptor) (CD antigen
Constituents: 50% Glycerol, 0.01M PBS, PH 7.4  >95%, Protein G purified  IgG  Polyclonal  Tumor necrosis factor receptor superfamily member 6 (Apo-1 antigen) (Apoptosis-mediating surface antigen FAS) (FASLG receptor) (CD antigen CD95), FAS, APT1 FAS1 TNFRSF6
Constituents: 50% Glycerol, 0.01M PBS, PH 7.4  >95%, Protein G purified  IgG  Polyclonal  Tumor necrosis factor receptor superfamily member 6 (Apo-1 antigen) (Apoptosis-mediating surface antigen FAS) (FASLG receptor) (CD antigen CD95), FAS, APT1 FAS1 TNFRSF6  Human

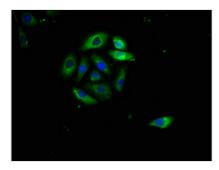


Immunohistochemistry of paraffin-embedded human placenta tissue using CSB-PA06354A0Rb at dilution of 1:100









Immunofluorescence staining of A549 cells with CSB-PA06354A0Rb at 1:400, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).