

### **Product datasheet for DM1212**

#### OriGene Technologies, Inc.

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# Fas Ligand (FASLG) Mouse Monoclonal Antibody [Clone ID: GM5F4]

### **Product data:**

**Product Type:** Primary Antibodies

Clone Name: GM5F4
Applications: FC, WB

**Recommended Dilution:** Flow Cytometry: 1.2 μg/10e6 cells.

ELISA: 1/200-1/400.

Reactivity: Human
Host: Mouse
Isotype: IgG1

Clonality: Monoclonal

**Immunogen:** Genetic immunisation with cDNA encoding Human FasL (extracellular domain)

**Specificity:** This antibody recognizes CD178 / Fas Ligand.

**Formulation:** Phosphate buffered saline, pH 7.2

State: Purified

State: Liquid purified Ig fraction

**Concentration:** lot specific

**Purification:** Affinity Chromatography on Protein G

**Conjugation:** Unconjugated

Storage: Store the antibody undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer.

Avoid repeated freezing and thawing.

**Stability:** Shelf life: one year from despatch.

Gene Name: Fas ligand

Database Link: Entrez Gene 356 Human

P48023





### Background:

The FAS ligand (FASL, CD95), a member of the tumor necrosis factor family, induces apoptosis in FAS-bearing cells (1). FASL is a type II membrane receptor with a soluble form that can be released into the extracellular fluid by proteolytic processing. Various cells express Fas, whereas FASL is expressed in activated splenocytes and thymocytes (2). In the immune system, Fas and FasL are involved in down-regulation of immune reactions as well as in T cell-mediated cytotoxicity. Malfunction of the Fas system causes lymphoproliferative disorders and accelerates autoimmune diseases, whereas its exacerbation may cause tissue destruction.

Synonyms:

FASLG, APT1LG1, FASL, TNFSF6, CD95L protein, APTL

# **Product images:**

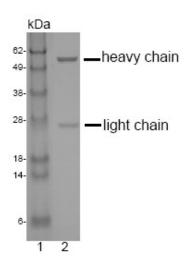


Fig.2: SDS-PAGE analysis of purified GM-5F4 monoclonalantibody. Lane 1: molecular weight marker, Lane 2: 2 ug ofpurified GM-5F4 antibody. Proteins were separated by SDS-PAGE and stained with RAPID Stain (TM) Reagent.

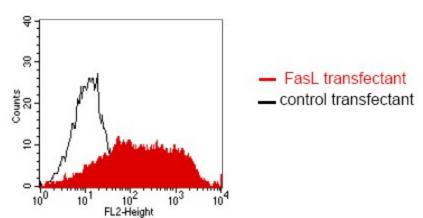


Fig.1: FACS analysis of BOSC23 cells using GM-5F4. BOSC23 cells were transiently transfected with an expres-sion vector encoding either FasL (red curve) or an irrelevantprotein (control transfectant: black curve). Binding of GM-5F4wasdetected with a PE-conjugated secondary antibody. A positivesignal was obtained only with FasL transfected cells.