bs-0687R

[Primary Antibody]

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DATASHEET -

Host: Rabbit Isotype: IgG

Clonality: Polyclonal

OAS1 Rabbit pAb

GeneID: 4938 SWISS: P00973

Target: OAS1

Immunogen: KLH conjugated synthetic peptide derived from human OAS1:

Purification: affinity purified by Protein A

Concentration: 1mg/ml

Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50%

Glycerol.

Shipped at 4°C. Store at -20°C for one year. Avoid repeated

freeze/thaw cycles.

Background: This protein is an enzyme included in the 2', 5' oligoadenylate

synthase family. This enzyme is induced by interferons and catalyzes the 2', 5' oligomers of adenosine in order to bind and activate RNase L. This enzyme family plays a significant role in the

inhibition of cellular protein synthesis and viral infection

resistance.

400-901-9800 Applications: WB (1:500-2000)

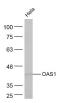
IHC-P (1:100-500) IHC-F (1:100-500) **IF** (1:100-500) Flow-Cyt (1ug/Test)

Reactivity: Human (predicted: Mouse,

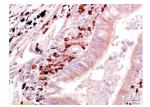
Predicted MW.: 40/46 kDa

Subcellular Location: Cytoplasm ,Nucleus

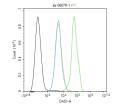
VALIDATION IMAGES



Sample: Hela(Human) Cell Lysate at 40 ug Primary: Anti-OAS1 (bs-0687R) at 1/300 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 40/46 kD Observed band size: 40 kD



Tissue/cell: human lung carcinoma; 4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer ($0.01\mbox{M}, \mbox{pH}\,6.0$), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Incubation: Anti-OAS1 Polyclonal Antibody, Unconjugated(bs-0687R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



Blank control (black line) :Hela. Primary Antibody (green line): Rabbit Anti-OAS1 antibody (bs-0687R) Dilution:1ug/Test; Secondary Antibody (white blue line): Goat anti-rabbit IgG-AF488 Dilution: 0.5ug/Test. Isotype control (orange line): Normal Rabbit IgG Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 90% ice-cold methanol for 20 min at -20°C, The cells were then incubated in 5%BSA to block nonspecific protein-protein interactions for 30 min at room temperature .Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.

- SELECTED CITATIONS -

• [IF=4.48] Xizhe Li. et al. Integrative bioinformatic analysis identified IFIT3 as a novel regulatory factor in psoriasis. J CELL BIOCHEM. 2022 Sep;: IHC; Human. 36169003