Product Datasheet

HGFR/c-MET Antibody (SJ19-05) NBP2-67412

Unit Size: 100 ul

Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.

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Updated 1/3/2023 v.20.1

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NBP2-67412

HGFR/c-MET Antibody (SJ19-05)

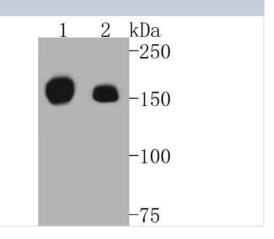
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Product Information		
Unit Size	100 ul	
Concentration	1 mg/ml	
Storage	Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.	
Clonality	Monoclonal	
Clone	SJ19-05	
Preservative	0.05% Sodium Azide	
Isotype	IgG	
Purity	Protein A purified	
Buffer	TBS (pH7.4), 0.05% BSA, 40% Glycerol	
Product Description		

Product Description	
Host	Rabbit
Gene Symbol	MET
Species	Human
	Synthetic peptide within C-terminal human HGFR/c-MET. (SwissProt: P08581 Human)

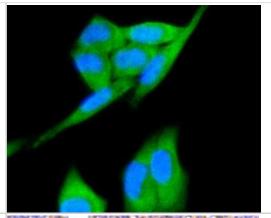
Product Application Details	
Applications	Western Blot, Flow Cytometry, Immunocytochemistry/ Immunofluorescence, Immunohistochemistry, Immunohistochemistry-Paraffin
Recommended Dilutions	Western Blot 1:1000-1:5000, Flow Cytometry 1:50-1:100, Immunohistochemistry, Immunocytochemistry/ Immunofluorescence 1:50-1:200, Immunohistochemistry-Paraffin 1:50-1:200

Images

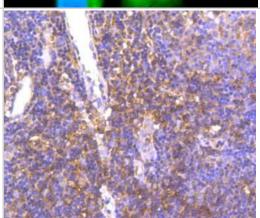
Western Blot: HGFR/c-MET Antibody (SJ19-05) [NBP2-67412] - Western blot analysis of HGFR/c-MET on different lysates. Proteins were transferred to a PVDF membrane and blocked with 5% BSA in PBS for 1 hour at room temperature. The primary antibody (1/500) was used in 5% BSA at room temperature for 2 hours. Goat Ant



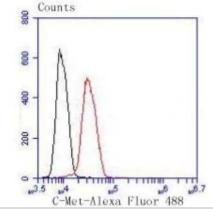
Immunocytochemistry/Immunofluorescence: HGFR/c-MET Antibody (SJ19-05) [NBP2-67412] - Staining Met in Hela cells (green). The nuclear counter stain is DAPI (blue). Cells were fixed in paraformaldehyde, permeabilised with 0.25% Triton X100/PBS.



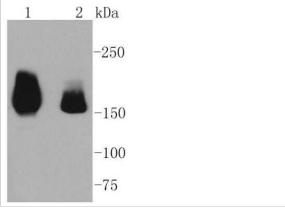
Immunohistochemistry-Paraffin: HGFR/c-MET Antibody (SJ19-05) [NBP2-67412] - Analysis of paraffin-embedded human tonsil tissue using anti-Met antibody. Counter stained with hematoxylin.



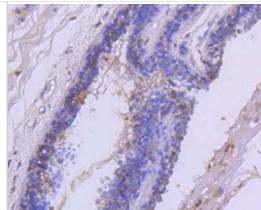
Flow Cytometry: HGFR/c-MET Antibody (SJ19-05) [NBP2-67412] - Flow cytometric analysis of HGFR/c-MET was done on Hela cells. The cells were fixed, permeabilized and stained with the primary antibody (1/50) (red). After incubation of the primary antibody at room temperature for an hour, the cells were stained with a Alexa Fluor 488-conjugated Goat anti-Rabbit IgG Secondary antibody at 1/1000 dilution for 30 minutes. Unlabeled sample was used as a control (cells without incubation with primary antibody; black).



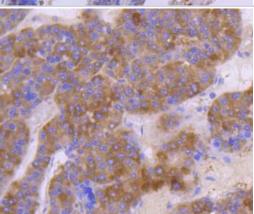
Western Blot: HGFR/c-MET Antibody (SJ19-05) [NBP2-67412] - Analysis of Met on different lysates using anti-Met antibody at 1/1,000 dilution. Positive control: Lane 1: Hela Lane 2: HepG2



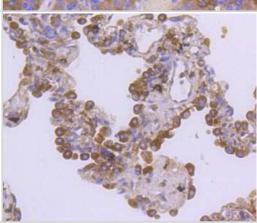
Immunohistochemistry-Paraffin: HGFR/c-MET Antibody (SJ19-05) [NBP2-67412] - Analysis of paraffin-embedded human breast carcinoma tissue using anti-Met antibody. Counter stained with hematoxylin.



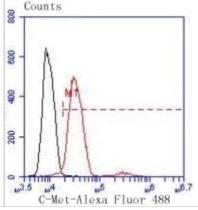
Immunohistochemistry-Paraffin: HGFR/c-MET Antibody (SJ19-05) [NBP2-67412] - Analysis of paraffin-embedded human liver cancer tissue using anti-Met antibody. Counter stained with hematoxylin.



Immunohistochemistry-Paraffin: HGFR/c-MET Antibody (SJ19-05) [NBP2-67412] - Analysis of paraffin-embedded human lung cancer tissue using anti-Met antibody. Counter stained with hematoxylin.



Flow Cytometry: HGFR/c-MET Antibody (SJ19-05) [NBP2-67412] - Analysis of Hela cells with Met antibody at 1/50 dilution (red) compared with an unlabelled control (cells without incubation with primary antibody; black). Alexa Fluor 488-conjugated goat anti rabbit IgG was used as the secondary antibody.





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Limitations

This product is for research use only and is not approved for use in humans or in clinical diagnosis. Primary Antibodies are guaranteed for 1 year from date of receipt.

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