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anti-MBTPS1 antibody (Internal Region)

3 Images



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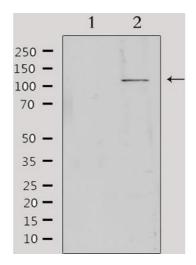
Target:

100 μL MBTPS1
MBTPS1
Internal Region
Human, Mouse, Rat
Rabbit
Polyclonal
This MBTPS1 antibody is un-conjugated
ELISA, Western Blotting (WB), Immunohistochemistry (IHC), Immunofluorescence (IF),
Immunocytochemistry (ICC)
A synthesized peptide derived from human MBTPS1, corresponding to a region within the
internal amino acids.
lgG
MBTPS1 Antibody detects endogenous levels of total MBTPS1.
Pig,Zebrafish,Bovine,Horse,Sheep,Rabbit,Dog,Chicken,Xenopus
The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling
Resin (Thermo Fisher Scientific).
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MBTPS1

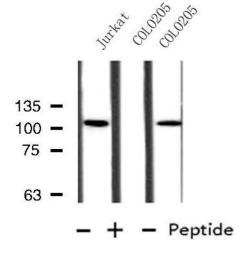
Target Details

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Alternative Name:	MBTPS1 (MBTPS1 Products)
Background:	Description: Serine protease that catalyzes the first step in the proteolytic activation of the
	sterol regulatory element-binding proteins (SREBPs). Other known substrates are BDNF,
	GNPTAB and ATF6. Cleaves after hydrophobic or small residues, provided that Arg or Lys is in
	position P4. Cleaves known substrates after Arg-Ser-Val-Leu (SERBP-2), Arg-His-Leu-Leu
	(ATF6), Arg-Gly-Leu-Thr (BDNF) and its own propeptide after Arg-Arg-Leu-Leu. Mediates the
	protein cleavage of GNPTAB into subunit alpha and beta, thereby participating in biogenesis of
	lysosomes.
	Gene: MBTPS1
Molecular Weight:	117 kDa
Gene ID:	8720
UniProt:	Q14703
Pathways:	ER-Nucleus Signaling
Application Details	
Application Notes:	WB 1:500-1:1000, IHC 1:50-1:200, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Concentration:	1 mg/mL
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 %
	glycerol.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which
	should be handled by trained staff only.
Storage:	-20 °C
Storage Comment:	Store at -20 °C. Stable for 12 months from date of receipt.
Expiry Date:	12 months



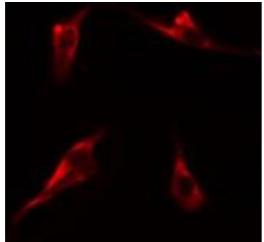
Western Blotting

Image 1. Western blot analysis of extracts from Hela, using MBTPS1 Antibody. Lane 1 was treated with the antigenspecific peptide.



Western Blotting

Image 2. Western blot analysis of extracts from Jurkat/COLO205 cells, using MBTPS1 antibody.



Immunofluorescence (fixed cells)

Image 3. ABIN6275227 staining Hela cells by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25¡ãC. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37¡ãC. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) antibody(Cat.# S0006), diluted at 1/600, was used as secondary antibod