# antibodies -online.com







# anti-Manic Fringe antibody (AA 28-321)



### Image



( )	1 /	-	rv	1 /	71	A
	1//	$\vdash$	1 \/	16		۱/۱
$\sim$	v	$\sim$	1 V	ı١	_	V١

Overview	
Quantity:	100 μL
Target:	Manic Fringe (MFNG)
Binding Specificity:	AA 28-321
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This Manic Fringe antibody is un-conjugated
Application:	ELISA, Immunohistochemistry (IHC)
Product Details	
Immunogen:	Recombinant Human Beta-1,3-N-acetylglucosaminyltransferase manic fringe protein (28-

Immunogen:	Recombinant Human Beta-1,3-N-acetylglucosaminyltransferase manic fringe protein (28-321AA)	
Isotype:	IgG	
Cross-Reactivity:	Human	
Purification:	Antigen Affinity Purified	

### Target Details

Target:	Manic Fringe (MFNG)	
Alternative Name:	MFNG (MFNG Products)	
Background: Background: Glycosyltransferase involved in the elongation of O-linked ligands to activ		

Notch signaling. Possesses fucose-specific beta-1,3-N-acetylglucosaminyltransferase activity. Aliases: 3-N-acetylglucosaminyltransferase manic fringe antibody, Beta-1 antibody, Beta-1,3-N-acetylglucosaminyltransferase manic fringe antibody, MFNG antibody, MFNG\_HUMAN antibody, O-fucosylpeptide 3-beta-N-acetylglucosaminyltransferase antibody

UniProt: 000587

Pathways: Notch Signaling

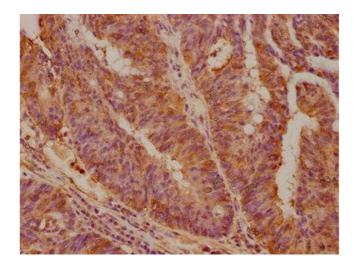
## **Application Details**

Application Notes:	Recommended dilution: IHC:1:20-1:200,
Restrictions:	For Research Use only

#### Handling

Format:	Liquid
Buffer:	Preservative: 0.03 % Proclin 300 Constituents: 50 % Glycerol, 0.01M PBS, pH 7.4
Preservative:	ProClin
Precaution of Use:	This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C,-80 °C
Storage Comment:	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.

#### **Images**



#### **Immunohistochemistry**

**Image 1.** IHC image of ABIN7145376 diluted at 1:100 and staining in paraffin-embedded human colon cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10 % normal goat serum 30 min at RT. Then primary antibody (1 % BSA) was incubated at 4 °C overnight. The primary is detected by a biotinylated secondary antibody and



visualized using an HRP conjugated SP system.