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# anti-GRM7 antibody (C-Term)





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Quantity:	100 μL
Target:	GRM7
Binding Specificity:	C-Term
Reactivity:	Human, Rat, Mouse
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This GRM7 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunofluorescence (IF), Immunocytochemistry (ICC)

## **Product Details**

Immunogen:	A synthesized peptide derived from human mGluR7, corresponding to a region within C-terminal amino acids.
Isotype:	IgG
Specificity:	MGluR7 Antibody detects endogenous levels of total mGluR7.
Predicted Reactivity:	Bovine,Horse,Dog
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink <sup>TM</sup> Coupling Resin (Thermo Fisher Scientific).

# Target Details

	GRM7	Target:
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# **Target Details**

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Alternative Name:	GRM7 (GRM7 Products)
Background:	Description: G-protein coupled receptor for glutamate. Ligand binding causes a conformation change that triggers signaling via guanine nucleotide-binding proteins (G proteins) and modulates the activity of down-stream effectors, such as adenylate cyclase. Signaling inhibits adenylate cyclase activity.  Gene: GRM7
Molecular Weight:	100kDa
Gene ID:	2917
UniProt:	Q14831
Pathways:	Sensory Perception of Sound, cAMP Metabolic Process, Feeding Behaviour
Application Details	
Application Notes:	WB: 1:500-1:3000, 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



# kDa 1 2 250— 150— 100— ——mGluR7 75— 50— 37— 25— 20— 15—



### **Immunohistochemistry**

**Image 1.** ABIN6266551 at 1/100 staining human brain tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue was then blocked and incubated with the antibody for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit antibody was used as the secondary.

### **Western Blotting**

**Image 2.** Western blot analysis on HuvEc cell lysate using mGluR7 Antibody,The lane on the left is treated with the antigen-specific peptide.

### Immunofluorescence (fixed cells)

**Image 3.** ABIN6266551 staining HuvEc 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°C. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary antibody.