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## anti-GABRg1 antibody (N-Term)

2 Images



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Overview	
Quantity:	100 μL
Target:	GABRg1
Binding Specificity:	N-Term
Reactivity:	Human, Rat, Mouse
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This GABRg1 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunofluorescence (IF), Immunocytochemistry (ICC)
Product Details	
Immunogen:	A synthesized peptide derived from human GABRG1, corresponding to a region within N-terminal amino acids.
Isotype:	IgG
Specificity:	GABRG1 Antibody detects endogenous levels of total GABRG1.
Predicted Reactivity:	Pig,Bovine,Horse,Sheep,Rabbit,Dog,Chicken
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink <sup>TM</sup> Coupling Resin (Thermo Fisher Scientific).
Target Details	
Target:	GABRg1

#### **Target Details**

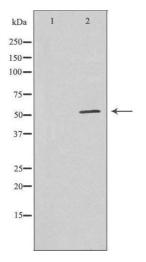
Alternative Name:	GABRG1 (GABRg1 Products)
Background:	Description: GABA, the major inhibitory neurotransmitter in the vertebrate brain, mediates
	neuronal inhibition by binding to the GABA/benzodiazepine receptor and opening an integral
	chloride channel.
	Gene: GABRG1
Molecular Weight:	54 kDa
Gene ID:	2565
UniProt:	Q8N1C3
Pathways:	Synaptic Membrane

### **Application Details**

Application Notes:	WB 1:500-1:1000, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000	
Restrictions:	For Research Use only	

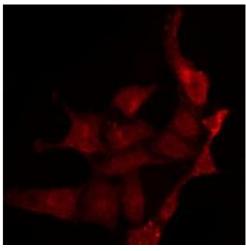
## Handling

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 RAW264.7 cells using GABRG1 antibody.



#### Immunofluorescence (fixed cells)

**Image 2.** ABIN6275140 staining LOVO 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) Ab, diluted at 1/600, was used as the secondary antibod