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## anti-GABRA4 antibody (AA 350-377)

**Images** 



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Quantity:	400 μL	
Target:	GABRA4	
Binding Specificity:	AA 350-377	
Reactivity:	Human	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This GABRA4 antibody is un-conjugated	
Application:	Western Blotting (WB), Flow Cytometry (FACS), Immunohistochemistry (Paraffin-embedded Sections) (IHC (p))	
Product Details		
Immunogen:	This GABRA4 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 350-377 amino acids from the Central region of human GABRA4.	
Isotype:	Ig Fraction	

### Target Details

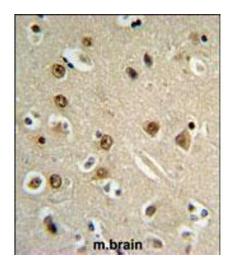
Purification:

Target:	GABRA4	
Alternative Name:	GABRA4 (GABRA4 Products)	
Background:	GABRA4 is the major inhibitory neurotransmitter in the mammalian brain where it acts at GABA-	
	A receptors, which are ligand-gated chloride channels. Chloride conductance of these channels	

This antibody is purified through a protein A column, followed by peptide affinity purification.

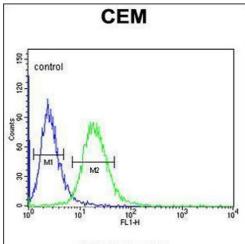
#### **Target Details**

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	can be modulated by agents such as benzodiazepines that bind to the GABA-A receptor.	
Molecular Weight:	62 kDa	
Gene ID:	2557	
UniProt:	P48169	
Pathways:	Synaptic Membrane	
Application Details		
Application Notes:	For WB starting dilution is: 1:1000	
	For IHC-P starting dilution is: 1:50~100	
	For FACS starting dilution is: 1:10~50	
Restrictions:	For Research Use only	
Handling		
Format:	Liquid	
Buffer:	Supplied in PBS with 0.09 % (W/V) sodium azide.	
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:	4 °C,-20 °C	
Storage Comment:	Store at 4°C for three months and -20°C, stable for up to one year. As with all antibodies care should be taken to avoid repeated freeze thaw cycles. Antibodies should not be exposed to prolonged high temperatures.	



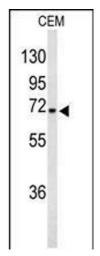
#### **Immunohistochemistry**

**Image 1.** GABRA4 Antibody IHC analysis in formalin fixed and paraffin embedded brain tissue followed by peroxidase conjugation of the secondary antibody and DAB staining.



#### **Flow Cytometry**

**Image 2.** Flow cytometric analysis of CEM cells (right histogram) compared to a negative control cell (left histogram).FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis.



#### **Western Blotting**

**Image 3.** Western blot analysis of GABRA4 Antibody in CEM cell line lysates (35ug/lane)