

Product datasheet

Anti-GABA A Receptor alpha 1 antibody ab32589

1 References 1 Image

Overview

<b>Product name</b>	Anti-GABA A Receptor alpha 1 antibody
<b>Description</b>	Rabbit polyclonal to GABA A Receptor alpha 1
<b>Host species</b>	Rabbit
<b>Tested applications</b>	<b>Suitable for:</b> WB
<b>Species reactivity</b>	<b>Reacts with:</b> Rat, Cow, Human
<b>Immunogen</b>	Fusion protein from the cytoplasmic loop of the alpha1-subunit of rat GABAA Receptor
<b>Positive control</b>	Rat, bovine, human and mouse brain extracts

Properties

<b>Form</b>	Liquid
<b>Storage instructions</b>	Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.
<b>Storage buffer</b>	Preservative: None Constituents: 50% Glycerol, 100µg/ml BSA, 150mM Sodium chloride, 10mM HEPES. pH 7.5
<b>Purity</b>	Immunogen affinity purified
<b>Purification notes</b>	Affinity purified
<b>Clonality</b>	Polyclonal
<b>Isotype</b>	IgG

Applications

Our [Abpromise guarantee](#) covers the use of **ab32589** in the following tested applications.

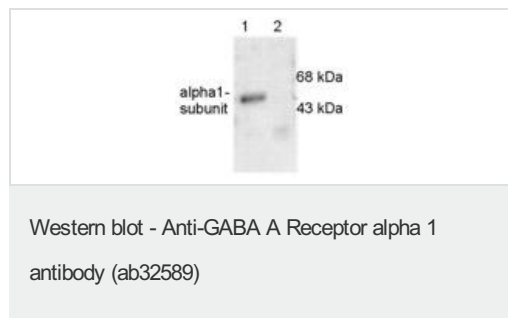
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB		1/1000. Detects a band of approximately 51 kDa (predicted molecular weight: 51 kDa).

Target

<b>Function</b>	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.
<b>Involvement in disease</b>	<p>Defects in GABRA1 are the cause of childhood absence epilepsy type 4 (ECA4) [MIM:611136]. A subtype of idiopathic generalized epilepsy characterized by onset at age 6-7 years, frequent absence seizures (several per day) and bilateral, synchronous, symmetric 3-Hz spike waves on EEG. During adolescence, tonic-clonic and myoclonic seizures may develop. Absence seizures may either remit or persist into adulthood.</p> <p>Defects in GABRA1 are the cause of juvenile myoclonic epilepsy type 5 (EJM5) [MIM:611136]. A subtype of idiopathic generalized epilepsy. Patients have afebrile seizures only, with onset in adolescence (rather than in childhood) and myoclonic jerks which usually occur after awakening and are triggered by sleep deprivation and fatigue.</p>
<b>Sequence similarities</b>	Belongs to the ligand-gated ion channel (TC 1.A.9) family. Gamma-aminobutyric acid receptor (TC 1.A.9.5) subfamily. GABRA1 sub-subfamily.
<b>Cellular localization</b>	Cell junction > synapse > postsynaptic cell membrane. Cell membrane.

## Images



**All lanes :** Anti-GABA A Receptor alpha 1 antibody (ab32589) at 1/1000 dilution

**Lane 1 :** Wildtype forebrain lysates 5-7 ug

**Lane 2 :** a1 knockout forebrain lysates 5-7 ug

**Predicted band size:** 51 kDa

**Observed band size:** 51 kDa

**Please note:** All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE"

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