# abcam

# Product datasheet

# Anti-alpha 1 Glycine Receptor/GLRA1 antibody ab228541

★★★★★ 1 Abreviews 1 References 3 图像

概述

免疫原

产品名称 Anti-alpha 1 Glycine Receptor/GLRA1抗体

描述 兔多克隆抗体to alpha 1 Glycine Receptor/GLRA1

**宿主** Rabbit

经测试应用 适用于: WB, IHC-P, ICC/IF

种属反应性 与反应: Rat, Human

预测可用于: Mouse, Cow, Zebrafish, Rhesus monkey \_\_\_\_\_\_

Recombinant fragment within Human alpha 1 Glycine Receptor/GLRA1 (internal sequence). The

exact sequence is proprietary.

Database link: P23415

阳性对照 WB: HeLa S3 whole cell lysate. IHC-P: OVCA xenograft tissue. ICC/IF: DIV9 rat E18 primary

cortical neuron cells.

常规说明

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

性能

形式 Liquid

**存放说明** Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

**存储溶液** pH: 7.00

Preservative: 0.01% Thimerosal (merthiolate)

Constituents: 1.21% Tris, 0.75% Glycine, 10% Glycerol (glycerin, glycerine)

纯**度** Immunogen affinity purified

 克隆
 多克隆

 同种型
 IgG

1

#### The Abpromise guarantee

#### Abpromise™承诺保证使用ab228541于以下的经测试应用

"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
WB	★ ☆ ☆ ☆ ☆ (1)	1/500 - 1/3000. Predicted molecular weight: 53 kDa.
IHC-P		1/100 - 1/1000.
ICC/IF		1/100 - 1/1000.

#### 靶标

功能 The glycine receptor is a neurotransmitter-gated ion channel. Binding of glycine to its receptor

increases the chloride conductance and thus produces hyperpolarization (inhibition of neuronal

firing).

疾病相关 Defects in GLRA1 are a cause of startle disease (STHE) [MIM:149400]; also known as hereditary

hyperekplexia or congenital stiff-person syndrome. STHE is a genetically heterogeneous

neurologic disorder characterized by muscular rigidity of central nervous system origin, particularly in the neonatal period, and by an exaggerated startle response to unexpected acoustic or tactile

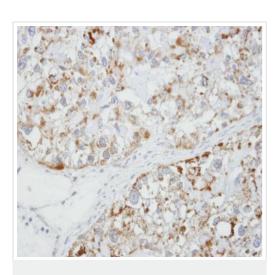
stimuli. Inheritance can be autosomal dominant or recessive.

**序列相似性** Belongs to the ligand-gated ion channel (TC 1.A.9) family. Glycine receptor (TC 1.A.9.3)

subfamily. GLRA1 sub-subfamily.

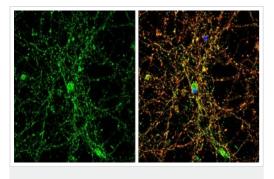
细胞定位 Cell junction > synapse > postsynaptic cell membrane. Cell membrane.

## 图片



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-alpha 1 Glycine
Receptor/GLRA1 antibody (ab228541)

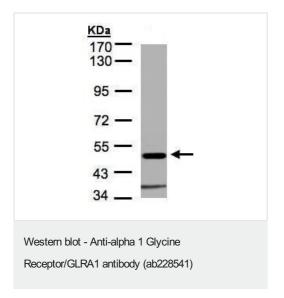
Paraffin-embedded OVCA (human) xenograft tissue stained for alpha 1 Glycine Receptor/GLRA1 using ab228541 at 1/100 dilution in immunohistochemical analysis.



Immunocytochemistry/ Immunofluorescence - Antialpha 1 Glycine Receptor/GLRA1 antibody (ab228541)

4% paraformaldehyde-fixed DIV9 rat E18 primary cortical neuron cells labeling alpha 1 Glycine Receptor/GLRA1 using ab228541 at 1/500 dilution.

The nuclear counterstain is DAPI (blue). beta Tubulin is labeled with an anti-beta Tubulin antibody (red).



Anti-alpha 1 Glycine Receptor/GLRA1 antibody (ab228541) at 1/1500 dilution + HeLa S3 whole cell lysate at 30  $\mu g$ 

Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 53 kDa

7.5% SDS-PAGE

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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