

## Datasheet

### GLRA1 purified MaxPab rabbit polyclonal antibody (D01P)

**Catalog Number:** H00002741-D01P

**Regulation Status:** For research use only (RUO)

**Product Description:** Rabbit polyclonal antibody raised against a full-length human GLRA1 protein.

**Immunogen:** GLRA1 (NP\_000162.2, 1 a.a. ~ 449 a.a) full-length human protein.

**Sequence:**

MYSFNTLRRLYLWETIVFFSLAASKEAEAARSAPKPMSP  
SDFLDKLMGRTSGYDARIRPNFKGPPVNVSCNIFINSF  
GSIAETMDYRVNIFLRQQWNDPRLAYNEYPDDSLDL  
DPSMLDSIWKPDLFFANEKGAFHEITTDNKLLRISRN  
GNVLYSIRITLTACPMDLKNFPMQVQTCIMQLESFGY  
TMNDLIFEWQEQGAVQVADGLTLPQFILKEEKDLRYCT  
KHYNTGKFTCIARFHLERQMGYYLIQMYIPSLILVLS  
WISFWINMDAAPARVGLGITTTLMTTQSSGSRASLPK  
VSYVKAIDIWMAVCLLFVFSALLEYAAVNFSRQHKEL  
LRFRRKRHHKEDEAGEGRFNFSAYGMGPACLQAKD  
GISVKGANNSNTTNPPPAPSKSPEEMRKLFIQRAKKID  
KISRIGFPM AFLIFNMFYWIIYKIVRREDVHNQ

**Host:** Rabbit

**Reactivity:** Human

**Applications:** WB-Tr

(See our web site product page for detailed applications information)

**Protocols:** See our web site at

<http://www.abnova.com/support/protocols.asp> or product page for detailed protocols

**Storage Buffer:** In 1x PBS, pH 7.4

**Storage Instruction:** Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

**Entrez GeneID:** 2741

**Gene Symbol:** GLRA1

**Gene Alias:** MGC138878, MGC138879, STHE

**Gene Summary:** The protein encoded by this gene is a subunit of a pentameric inhibitory glycine receptor. The receptor mediates postsynaptic inhibition in the central nervous system. Defects in this gene are a cause of startle disease (STHE), also known as hereditary hyperekplexia or congenital stiff-person syndrome. Two transcript variants encoding different isoforms have been found for this gene]