

CHRNE Antibody (Center)

Purified Rabbit Polyclonal Antibody (Pab)

Catalog # AP22287c

Product Information

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|--------------------------|---|
| Application | WB, FC, E |
| Primary Accession | Q04844 |
| Other Accession | P20782 , P09660 |
| Reactivity | Human, Rat, Mouse |
| Predicted | Mouse, Rat |
| Host | Rabbit |
| Clonality | polyclonal |
| Isotype | Rabbit IgG |
| Clone Names | RB56765 |
| Calculated MW | 54697 |

Additional Information

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|---------------------------|--|
| Gene ID | 1145 |
| Other Names | Acetylcholine receptor subunit epsilon, CHRNE, ACHRE |
| Target/Specificity | This CHRNE antibody is generated from a rabbit immunized with a KLH conjugated synthetic peptide between 409-443 amino acids from the Central region of human CHRNE. |
| Dilution | WB~~1:2000 FC~~1:25 E~~Use at an assay dependent concentration. |
| Format | Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification. |
| Storage | Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles. |
| Precautions | CHRNE Antibody (Center) is for research use only and not for use in diagnostic or therapeutic procedures. |

Protein Information

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|-----------------|--|
| Name | CHRNE (HGNC:1966) |
| Synonyms | ACHRE |
| Function | After binding acetylcholine, the AChR responds by an extensive change in conformation that affects all subunits and leads to opening of an |

ion-conducting channel across the plasma membrane.

Cellular Location

Postsynaptic cell membrane; Multi-pass membrane protein. Cell membrane; Multi-pass membrane protein

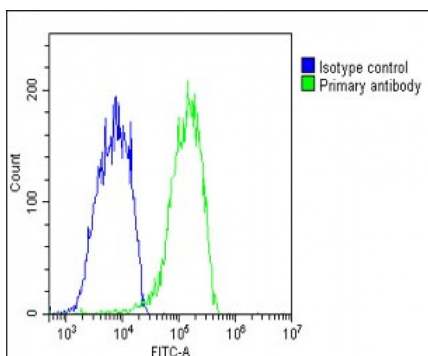
Background

After binding acetylcholine, the AChR responds by an extensive change in conformation that affects all subunits and leads to opening of an ion-conducting channel across the plasma membrane.

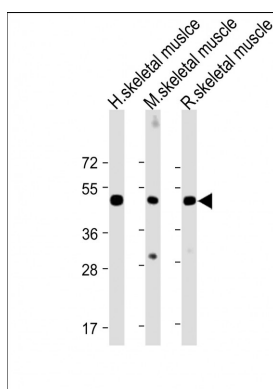
References

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Mural R.J.,et al.Submitted (SEP-2005) to the EMBL/GenBank/DDBJ databases.
Gomez C.M.,et al.Neurology 45:982-985(1995).
Ohno K.,et al.Proc. Natl. Acad. Sci. U.S.A. 92:758-762(1995).

Images



Overlay histogram showing HepG2 cells stained with AP22287c(green line). The cells were fixed with 2% paraformaldehyde (10 min). The cells were then incubated in 2% bovine serum albumin to block non-specific protein-protein interactions followed by the antibody (AP22287c, 1:25 dilution) for 60 min at 37°C. The secondary antibody used was Goat-Anti-Rabbit IgG, DyLight® 488 Conjugated Highly Cross-Adsorbed(1583138) at 1/200 dilution for 40 min at 37°C. Isotype control antibody (blue line) was rabbit IgG1 (1µg/1x10⁶ cells) used under the same conditions. Acquisition of >10, 000 events was performed.



All lanes : Anti-CHRNE Antibody (Center) at 1:2000 dilution Lane 1: Human skeletal muscle lysate Lane 2: Mouse skeletal muscle lysate Lane 3: Rat skeletal muscle lysate Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution. Predicted band size : 55 kDa Blocking/Dilution buffer: 5% NFDM/TBST.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.