

CHRND Rabbit pAb

货号: B20089

产品信息

反应	Human,Mouse
宿主	Rabbit
克隆性	Polyclonal
预测反应	
应用	WB
推荐浓度	WB: 1:500 - 1:2000
理论分子量	57kDa/58kDa
实测分子量	59kDa
形式	Liquid
保存条件	Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.01% thiomersal,50% glycerol,pH7.3.
偶联物	Unconjugated
阳性对照	A-431,293T,HeLa,Mouse lung
细胞定位	Cell junction,Cell membrane,Multi-pass membrane protein,postsynaptic cell membrane,synapse
纯化	Affinity purification

抗原信息

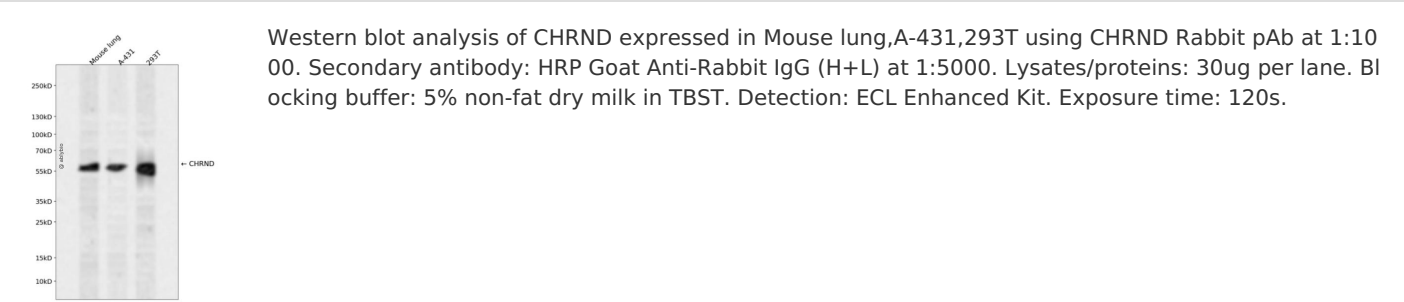
抗原信息	Recombinant fusion protein containing a sequence corresponding to amino acids 22-245 of human CHRND (NP_000742.1).
序列	LNEEERLIRHLFQEKGYNKELRPVAHKEESVDVALALTSLNLISLKEVEETLTTNVWIEHGWTDNRLKWNAEEFGNISVLRLP PDMVWLPEIVLENNNDGSFQISYSCNVLVYHYGFVYWLPPAIFRSSCPISVTYFPFDWQNCSLKFSSLKYTAKEITLSLKQD AKENRTYPVEWIIIDPEGFTENGWEIVHRPARVNVDPRAPLDSRQDITFYLIIRRK

靶点信息

研究背景	The acetylcholine receptor of muscle has 5 subunits of 4 different types: 2 alpha and 1 each of beta, gamma and delta subunits. After acetylcholine binding, the receptor undergoes an extensive conformation change that affects all subunits and leads to opening of an ion-conducting channel across the plasma membrane. Defects in this gene are a cause of multiple pterygium syndrome lethal type (MUPSL), congenital myasthenic syndrome slow-channel type (SCCMS), and congenital myasthenic syndrome fast-channel type (FCCMS). Several transcript variants encoding different isoforms have been found for this gene.
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基因ID	1144
基因名	CHRND
Swiss	Q07001
别名	CHRND;ACHRD;CMS2A;CMS3A;CMS3B;CMS3C;FCCMS;SCCMS

产品验证



实验步骤

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