

# NEU1 Rabbit pAb

货号: B16794

产品信息

反应	Human,Mouse,Rat
宿主	Rabbit
克隆性	Polyclonal
预测反应	
应用	WB IHC
推荐浓度	<b>WB:</b> 1:500 - 1:1000 <b>IHC:</b> 1:50 - 1:200
理论分子量	45kDa
实测分子量	45KDa
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	BxPC-3,Mouse kidney,Mouse pancreas,Rat liver
细胞定位	Cell membrane,Cytoplasmic vesicle,Lumenal side,Lysosome lumen,Lysosome membrane,Peripheral mem brane protein
纯化	Affinity purification

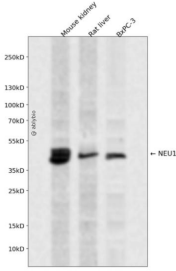
抗原信息

抗原信息	Recombinant fusion protein containing a sequence corresponding to amino acids 210-415 of human NEU 1 (NP_000425.1).
序列	PRKGRLIVCGHGTLERDGVFCLLSDDHGASWRYGSGVSGIPYGQPKQENDFNPDECQPYELPDGSVVINARNQNNYHC HCRIVLRSYDACDTLRPRDVTDFPELVDPVVAAGAVVTSSGIVFFSNPAHPEFRVNLTLRWSFSNGTSWRKETVQLWPGP SGYSSLATLEGSMDGEEQAPQLYVLYEKGRNHYTESISVAKISVYGTL

靶点信息

研究背景	The protein encoded by this gene is a lysosomal enzyme that cleaves terminal sialic acid residues from s substrates such as glycoproteins and glycolipids. In the lysosome, this enzyme is part of a heterotrimeric c omplex together with beta-galactosidase and cathepsin A (the latter is also referred to as 'protective prot ein'). Mutations in this gene can lead to sialidosis, a lysosomal storage disease that can be type 1 (cherry red spot-myoclonus syndrome or normosomatic type), which is late-onset, or type 2 (the dysmorphic type ), which occurs at an earlier age with increased severity.
基因ID	4758
基因名	NEU1
Swiss	Q99519
别名	NEU1;NANH;NEU;SIAL1

产品验证



Western blot analysis of NEU1 expressed in Mouse kidney,Rat liver,BxPC-3 using NEU1 Rabbit pAb at 1: 1000. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) at 1:5000. Lysates/proteins: 30ug per lane. Blocking buffer: 5% non-fat dry milk in TBST. Detection: ECL Enhanced Kit. Exposure time: 120s.

实验步骤

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