

Datasheet for ABIN5692834 anti-CNR1 antibody (AA 1-75)





Overview

Quantity:	100 μg
Target:	CNR1
Binding Specificity:	AA 1-75
Reactivity:	Human, Rat, Mouse
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This CNR1 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Flow Cytometry (FACS), Immunocytochemistry (ICC)

Product Details

Purpose:

Immunogen:

Isotype:	IgG
Cross-Reactivity (Details):	No cross-reactivity with other proteins.
Characteristics:	Anti-Cannabinoid Receptor I/CNR1 Antibody Picoband® (ABIN5692834). Tested in ELISA, Flow
	Cytometry, IHC, ICC, WB applications. This antibody reacts with Human, Mouse, Rat. The brand
	Picoband indicates this is a premium antibody that guarantees superior quality, high affinity,
	and strong signals with minimal background in Western blot applications. Only our best-
	performing antibodies are designated as Picoband, ensuring unmatched performance.

E. coli-derived human Cannabinoid Receptor I recombinant protein (Position: M1-Q75).

Anti-Cannabinoid Receptor I/CNR1 Antibody Picoband®

Target Details

Target:	CNR1		
Alternative Name:	CNR1 (CNR1 Products)		
Background:	Synonyms: Cannabinoid receptor 1, CB-R, CB1, CANN6, CNR1, CNR		
	Tissue Specificity: Widely expressed, with highest levels in fetal and adult brain. Expression		
	levels of isoform 2 and isoform 3 are much lower than those of isoform 1.		
	Background: The cannabinoid receptor type 1, often abbreviated as CB1, is a G protein-coupled		
	cannabinoid receptor located primarily in the central and peripheral nervous system. This gene		
	encodes one of two cannabinoid receptors. The cannabinoids, principally delta-9-		
	tetrahydrocannabinol and synthetic analogs, are psychoactive ingredients of marijuana. The		
	cannabinoid receptors are members of the guanine-nucleotide-binding protein (G-protein)		
	coupled receptor family, which inhibit adenylate cyclase activity in a dose-dependent,		
	stereoselective and pertussis toxin-sensitive manner. The two receptors have been found to be		
	involved in the cannabinoid-induced CNS effects (including alterations in mood and cognition)		
	experienced by users of marijuana. Multiple transcript variants encoding two different protein		
	isoforms have been described for this gene.		
Molecular Weight:	60 kDa		
Gene ID:	1268		
UniProt:	P21554		
Pathways:	Feeding Behaviour		
Application Details			
Application Notes:	Western blot, 0.1-0.5 μg/mL		
	Immunohistochemistry (Paraffin-embedded Section), 0.5-1 μg/mL		
	Immunohistochemistry (Frozen Section), 0.5-1 μg/mL		
	Immunocytochemistry, 0.5-1 μg/mL		
	Flow Cytometry (Fixed), 1-3 µg/1x10 ⁶ cells		
	ELISA, 0.1-0.5 μg/mL		
	1. "Entrez Gene: CNR1 cannabinoid receptor 1 (brain)". 2. Russo, P., Strazzullo, P., Cappuccio, F.		
	P., Tregouet, D. A., Lauria, F., Loguercio, M., Barba, G., Versiero, M., Siani, A. Genetic variations at		
	the endocannabinoid type 1 receptor gene (CNR1) are associated with obesity phenotypes in		
	men. J. Clin. Endocr. Metab. 92: 2382-2386, 2007.		
Restrictions:	For Research Use only		

Handling

Format:	Lyophilized
Reconstitution:	Add 0.2 mL of distilled water will yield a concentration of 500 $\mu g/mL$.
Concentration:	500 μg/mL
Buffer:	Each vial contains 4 mg Trehalose, 0.9 mg NaCl, 0.2 mg Na $_2$ HPO $_4$, 0.05 mg NaN $_3$.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	4 °C,-20 °C
Storage Comment:	Store at -20°C for one year from date of receipt. After reconstitution, at 4°C for one month. It can also be aliquotted and stored frozen at -20°C for six months. Avoid repeated freeze-thaw cycles.

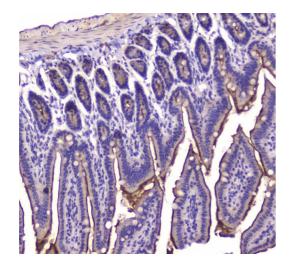
Images

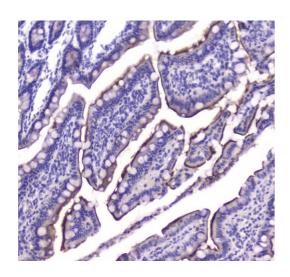
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Western Blotting

Image 1. Western blot analysis of Cannabinoid Receptor I using anti-Cannabinoid Receptor I antibody. Electrophoresis was performed on a 5-20% SDS-PAGE gel at 70V (Stacking gel) / 90V (Resolving gel) for 2-3 hours. The sample well of each Lane was loaded with 50ug of sample under reducing conditions. Lane 1: human 22RV1 whole cell lysates. After Electrophoresis, proteins were transferred to Nitrocellulose membrane at 150mA for 50-90 minutes. Blocked the membrane with 5% Non-fat Milk/ TBS for 1.5 hour at RT. The membrane was incubated with rabbit anti-Cannabinoid Receptor I antigen affinity purified polyclonal antibody (Catalog #) at 0.5 µg/mL overnight at 4°C, then washed with TBS-0.1%Tween 3 times with 5 minutes each and probed with a goat anti-rabbit IgG-HRP secondary antibody at a dilution of 1:10000 for 1.5 hour at RT. The signal is developed using an Enhanced Chemiluminescent detection (ECL) kit (Catalog # EK1002) with Tanon 5200 system. A specific band was detected for Cannabinoid





Receptor I at approximately 60KD. The expected band size for Cannabinoid Receptor I is at 53KD.

Immunohistochemistry

Image 2. IHC analysis of Cannabinoid Receptor I using anti-Cannabinoid Receptor I antibody . Cannabinoid Receptor I was detected in paraffin-embedded section of mouse small intestine tissue. Heat mediated antigen retrieval was performed in citrate buffer (pH6, epitope retrieval solution) for 20 mins. The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 1μg/ml rabbit anti-Cannabinoid Receptor I Antibody overnight at 4°C. Biotinylated goat anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Strepavidin-Biotin-Complex (SABC)(Catalog # SA1022) with DAB as the chromogen.

Immunohistochemistry

Image 3. IHC analysis of Cannabinoid Receptor I using anti-Cannabinoid Receptor I antibody. Cannabinoid Receptor I was detected in paraffin-embedded section of rat small intestine tissue. Heat mediated antigen retrieval was performed in citrate buffer (pH6, epitope retrieval solution) for 20 mins. The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 1μg/ml rabbit anti-Cannabinoid Receptor I Antibody overnight at 4°C. Biotinylated goat anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Strepavidin-Biotin-Complex (SABC)(Catalog # SA1022) with DAB as the chromogen.