## KCNQ2 Rabbit Polyclonal Antibody

## CAB1917



Product Information	Protein Background
Size:	The M channel is a slowly activating and deactivating potassium channel that plays a critica
20uL, 50uL, 100uL, 200uL	role in the regulation of neuronal excitability. The M channel is formed by the association of the protein encoded by this gene and a related protein encoded by the KCNQ3 gene, both integral membrane proteins. M channel currents are inhibited by M1 muscarinic acetylcholine receptors and activated by retigabine, a novel anti-convulsant drug. Defects in this gene are a
Observed MW:	
96kDa	cause of benign familial neonatal convulsions type 1 (BFNC), also known as epilepsy, benigr neonatal type 1 (EBN1). At least five transcript variants encoding five different isoforms have
Calculated MW:	been found for this gene.
44kDa/92kDa/93kDa/94kDa/ 95kDa	Immunogen information
Applications:	Gene ID: 3785
WB	
Reactivity:	Uniprot O43526
Human, Mouse, Rat	
	<b>Synonyms:</b> KCNQ2; BFNC; EBN; EBN1; ENB1; HNSPC; KCNA11; KV7.2
Antibody Information	
Recommended dilutions:	
WB 1:500 - 1:2000	Immunogen:
	Recombinant fusion protein containing a sequence corresponding
<b>Source:</b> Rabbit	to amino acids 254-393 of human KCNQ2 (NP_742107.1).
	Storage:
lsotype:	Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.02%
igu	source after $a_2$ and $a_3$ by $a_6$ gryceror, $\mu \pi r$ .s.

**Purification:** Affinity purification

## **Product Images**



Western blot analysis of extracts of Raji cells, using KCNQ2 antibody (CAB1917) at 1:3000 dilution. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (CABS014) at 1:10000 dilution. Lysates/proteins: 25ug per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (CABM00020). Exposure time: 90s.