



DCAMKL2 rabbit pAb

Cat#: orb765025 (Manual)

For research use only. Not intended for diagnostic use.

Product Name DCAMKL2 rabbit pAb

Host species Rabbit

Applications WB;IF;ELISA

Species Cross-Reactivity Human; Mouse

Recommended dilutions Western Blot: 1/500 - 1/2000. Immunofluorescence: 1/200 - 1/1000. ELISA:

1/10000. Not yet tested in other applications.

Immunogen The antiserum was produced against synthesized peptide derived from

human DCLK2. AÁ range:1-50

Specificity DCAMKL2 Polyclonal Antibody detects endogenous levels of DCAMKL2

protein.

Formulation Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium

azide..

Storage Store at -20°C. Avoid repeated freeze-thaw cycles.

Protein Name Serine/threonine-protein kinase DCLK2

Gene Name DCLK2

Cellular localization Cytoplasm, cytoskeleton. Colocalizes with microtubules. .

Purification The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Clonality Polyclonal





Concentration 1 mg/ml

Observed band 83kD

Human Gene ID 166614

Human Swiss-Prot Number Q8N568

DCLK2; DCAMKL2; DCDC3B; DCK2; Serine/threonine-protein kinase DCLK2; CaMK-like CREB regulatory kinase 2; CL2; CLICK-II; CLICK2; **Alternative Names**

Doublecortin domain-containing protein 3B; Doublecortin-like and CAM

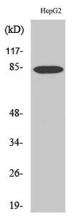
kinase-like 2; Doublecortin-like kinase 2

Background This gene encodes a member of the protein kinase superfamily and the

doublecortin family. The protein encoded by this gene contains two Nterminal doublecortin domains, which bind microtubules and regulate microtubule polymerization, a C-terminal serine/threonine protein kinase domain, which shows substantial homology to Ca2+/calmodulin-dependent protein kinase, and a serine/proline-rich domain in between the doublecortin and the protein kinase domains, which mediates multiple protein-protein interactions. The microtubule-polymerizing activity of the encoded protein is independent of its protein kinase activity. Mouse studies show that the DCX gene, another family member, and this gene share function in the

establishment of hippocampal organization and that their absence results in a severe epileptic phenotype and lethality, as described in human patients with

lissencephaly. Multiple alterna

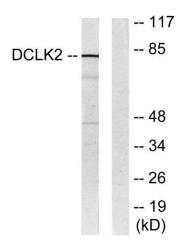


Western Blot analysis of various cells using DCAMKL2 Polyclonal Antibody





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Western blot analysis of lysates from HepG2 cells, using DCLK2 Antibody. The lane on the right is blocked with the synthesized peptide.