

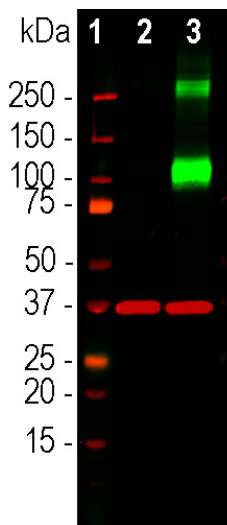
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HGNC Name: ADCY3
UniProt: P21932
RRID: AB_2744501
Immunogen: C-terminal peptide of rat ACIII, PAAFPNGSSVTLPHQVVDNP with a Cys added to the N-terminus to allow coupling to KLH.
Format: Purified antibody at 1mg/mL in 50% PBS, 50% glycerol plus 5mM NaH₂PO₄
Storage: Store at 4°C for short term, for longer term at -20°C
Recommended dilutions:
WB: 1:1,000-1:2,000. IF/ICC and IHC: 1:1,000

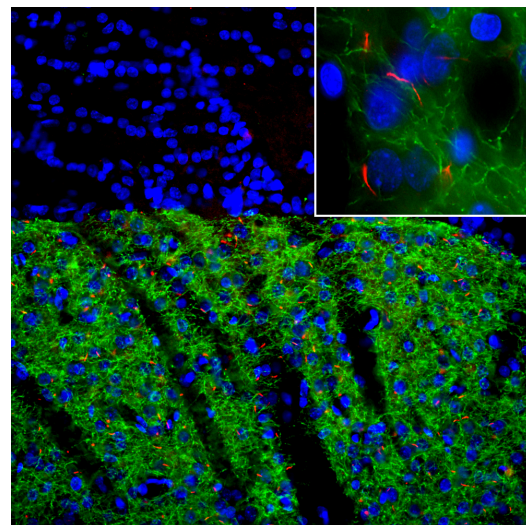
References:

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2. Louvi A and Grove EA. Cilia in the CNS: the quiet organelle claims center stage. *Neuron* 69:1046-60 (2011).
3. Singla V, Reiter JF. The primary cilium as the cell's antenna: signaling at a sensory organelle. *Science* 313:629-33 (2006).
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6. Gomez-Gamboa A, et al. Primary cilia in the developing and mature brain. *Neuron* 82:511-21 (2014).
7. Guadiana SM, et al. Arborization of Dendrites by developing neocortical neurons is dependent on primary cilia and Type 3 adenylyl cyclase. *J. Neurosci.* 33:2626-38 (2013).
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Applications	Host	Isotype	Molecular Wt.	Species Cross-Reactivity
WB, IF/ICC, IHC	Mouse	IgG1	~120kDa and above	Hu, Rt, Ms



Western blot analysis of HEK293 cell lysates using mouse mAb to ACIII, MCA-1A12, dilution 1:1,000, in green: [1] protein standard, [7] non-transfected HEK293 cells, and [8] HEK293 cells transfected with a vector encoding Myc-DDK tagged full length human adenylate cyclase III (ACIII). The strong band at about 130kDa in transfected cells demonstrates overexpression of the human ACIII protein, and bands over 250kDa presumably correspond to either heavily glycosylated form of ACIII or multimeric forms of the protein. The same blot was simultaneously probed with rabbit pAb to GAPDH, [RPCA-GAPDH](#), dilution 1:20,000 in red, which reveals the single band at ~37kDa seen in both transfected and non-transfected cells.



Immunofluorescence of caudate/putamen region of rat brain stained with mouse mAb to ACIII, MCA-1A12, in red, and chicken antibody to tyrosine hydroxylase, in green. The Blue is Hoechst stain revealing nuclei. The ACIII antibody reveals neuronal cilia while the tyrosine hydroxylase antibody reveals the axons of chatecholaminergic neurons.

Background:

Trimeric G-proteins are a large and variable family of membrane receptors. On binding their specific ligand they activate specific members of the family of trimeric G-proteins which in turn activate other signalling enzymes. Adenylate cyclases are one of these downstream enzyme families which are activated by the GTP bound Gαs subunits of trimeric G-proteins. Adenylate cyclases are responsible for the production of the important "second messenger" signaling molecule cyclic-AMP which in turn activates the cAMP dependent protein kinase. This kinase when activated phosphorylates numerous substrate molecules on serine or threonine residues and so alters their activity. There are several different adenylate cyclase genes and protein products with each have distinctly different distribution patterns in cells and tissues. The type III adenylate cyclase enzyme is specifically localized in the membranes surrounding neuronal cilia, and is activated by specific G-protein coupled receptors also located in cilia (1-5). Neuronal cilia express a variety of other receptors types and mediators of other signaling pathways and appear to function as a unique and complex neuronal sensory structure (1-5). For examples, the somatostatin 3 receptor, neuropeptide Y 2 receptor and melanin concentrating hormone receptor 1 are localized in neuronal cilia and the sonic hedgehog and Wnt signalling pathway act on neurons primarily through neuronal cilia (6). This antibody is an excellent marker of neuronal cilia in the brain and in cells in tissue culture and works in the same way as our rabbit polyclonal made against the same peptide (7). The antibody was recently utilized in a very high profile publication in the journal *Cell* (8).

The MCA-1A12 antibody was made against the extreme C-terminal peptide of rat ACIII, PAAFPNGSSVTLPHQVVDNP, amino acids 1125-1144 of the Genbank entry [NP_570135.2](#). A cysteine residue was added to the N-terminus to allow coupling to MBS-activated keyhole limpet hemocyanin. The antibody works on mouse cells which express the same peptide and also on human cells, presumably because the corresponding peptide in the human AC3 sequence is the closely related peptide LATFPNGPSVTLPHQVVDNS. The antibody works well to identify neuronal cilia on both human and rodent cells by IF and ICC but is not recommended for IHC. We have also generated rabbit and chicken polyclonal antibodies to the same ACIII peptide, [RPCA-ACIII](#) and [CPCA-ACIII](#).

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Abbreviation Key:

mAb—Monoclonal Antibody **pAb**—Polyclonal Antibody **WB**—Western Blot **IF**—Immunofluorescence **ICC**—Immunocytochemistry
IHC—Immunohistochemistry **E**—ELISA **Hu**—Human **Mo**—Monkey **Do**—Dog **Rt**—Rat **Ms**—Mouse **Co**—Cow **Pi**—Pig **Ho**—Horse **Ch**—Chicken
Dr—*D. rerio* **Dm**—*D. melanogaster* **Sm**—*S. mutans* **Ce**—*C. elegans* **Sc**—*S. cerevisiae* **Sa**—*S. aureus* **Ec**—*E. coli*.

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IHC—Immunohistochemistry E—ELISA Hu—Human Mo—Monkey Do—Dog Rt—Rat Ms—Mouse Co—Cow Pi—Pig Ho—Horse Ch—Chicken
Dr—D. rerio Dm—D. melanogaster Sm—S. mutans Ce—C. elegans Sc—S. cerevisiae Sa—S. aureus Ec—E. coli.*