

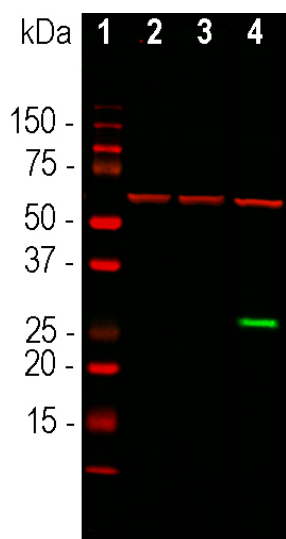
Ordering Information
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HGNC Name: NA
UniProt: Q6YGZ0
RRID: AB_2572315
Immunogen: The prot-r-AcGFP recombinant protein purified from *E. coli*. The epitope is in the N-terminal 18 amino acids of the protein, the peptide MVSKGAELFTGIVPLIE, which is found in the Clontech and other GFP vectors
Format: Purified antibody at 1mg/mL in 50% PBS, 50% glycerol plus 5mM NaN₃
Storage: Stable at 4°C for one year, for longer term store at -20°C
Recommended dilutions:
WB: 1:1,000, IF/ICC 1:1,000

References:

1. Shimomura O, Johnson FH, Saiga Y. Extraction, purification and properties of aequorin, a bioluminescent protein from the luminous hydromedusa, *Aequorea*. *J. Cell. Comp. Physiol.* 3:223-39 (1962).
2. Shimomura, O. Structure of the chromophore of *Aequorea* green fluorescent protein. *FEBS Lett.* 104:220-2 (1979).
3. Prasher DC, et al. Primary structure of the *Aequorea victoria* green-fluorescent protein. *Gene* 111:229-33 (1992).
4. Cody CW, et al. Chemical structure of the hexapeptide chromophore of the *Aequorea* green-fluorescent protein. *Biochem.* 32:1212-8 (1993).
5. Chalfie M, et al. Green Fluorescent protein as a marker for gene expression. *Science* 263:802-5 (1994).
6. Heim R, Prasher DC, Tsien RY. Wavelength mutations and post-translational autooxidation of green fluorescent protein. *PNAS* 91:12501-04 (1994).
7. Ormo M, et al. Crystal structure of the *Aequorea victoria* green fluorescent protein. *Science* 273:1392-95 (1996).
8. Tsien RY. The green fluorescent protein. *Annu. Rev. Biochem.* 67:509-44 (1998).
9. Zacharias DA, Violin JD, Newton AC, Tsien RY. Partitioning of lipid-modified monomeric GFPs into membrane microdomains of live cells. *Science* 296:913-6 (2002).
10. Gurskaya NG, et al. A colourless green fluorescent protein homologue from the non-fluorescent hydromedusa *Aequorea coerulescens* and its fluorescent mutants. *Biochem. J.* 373:403-8 (2003).

Applications	Host	Isotype	Molecular Wt.	Species Cross-Reactivity
WB, ICC/IF, IHC	Mouse	IgM	~27kDa	NA

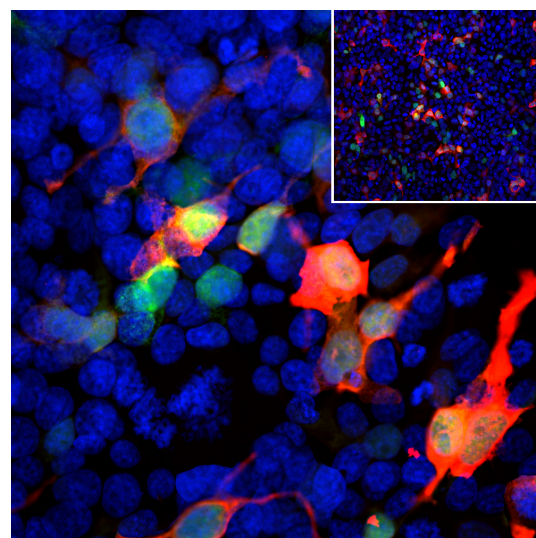


Western blot analysis of transfected and control HEK293 cell lysates using mouse mAb to GFP, MCA-1F1, in green, dilution 1:1,000: [1] protein standard, [2] Control, non-transfected cells, [3] cells transfected with an mCherry red fluorescent protein construct and [4] cells transfected with GFP construct. The strong green band at ~27kDa corresponds to GFP protein detected only in cells transfected with GFP construct, the antibody does not bind to mCherry. The same blot was simultaneously probed with chicken pAb to HSP60, CPCA-HSP60, dilution 1:10,000, in red. The single band at 60kDa represents the HSP60 protein expressed in all preparations.

Background:

The **green fluorescent protein** (GFP) is a 27kDa protein isolated originally from the jellyfish *Aequorea victoria*. It has an endogenous fluorochrome activity with excitation maximum at 395nm and emission maximum at 509nm, which is similar to that of fluorescein (1,2). The GFP gene was sequenced and the origin of the fluorochrome by autocatalytic activity of certain amino acids was discovered (3,4). Much interest in GFP was generated when it was shown that fluorescence develops rapidly when the protein is expressed and requires only molecular oxygen and no other cofactors. As a result GFP can be expressed in fluorescent form in essentially any prokaryotic or eukaryotic cell (5). GFP has been engineered to produce a vast number of variously colored mutants including blue, cyan and yellow protein derivatives, BFP, CFP and YFP (6-9). GFP and other fluorescent proteins derived from other Cnidarians (jellyfish, coral and medusa) are widely used as tracers in transfection and transgenic experiments to monitor gene expression and protein localization *in vivo* and *in vitro*. The crystal structure of GFP was determined (7) which allowed amino acid modifications to improve spectral properties and prevent multimerization (8,9). The discovery and use of GFP was the basis of the **2008 Nobel prize in chemistry**, specifically "for the discovery and development of the green fluorescent protein, GFP".

The MCA-1F1 antibody was made against a recombinant GFP construct originating from an *Aequorea* species which was engineered to improve spectral properties and prevent oligomerization (10). This form of GFP, referred to as AcGFP, is 94% identical to the eGFP developed by Tsien and coworkers and is the form of GFP inserted in the **Clontech/Takara expression vectors**. We epitope mapped this antibody to the N-terminal 18 amino acids of the protein, the peptide MVSKGAELFTGIVPLIE, and showed that the antibody binds the similar N-terminal peptide of eGFP. For detailed sequence information see [here](#). We also supply the immunogen, **PROT-AcGFP**. The antibody can be used to verify the expression, size and stability of both AcGFP and eGFP fusion proteins in western blotting experiments and to amplify GFP signals in tissues of transgenic animals. We also supply another mouse monoclonal antibody which has a different isotype and rabbit, chicken, goat polyclonal antibodies to this protein, **MCA-3B11**, **RPCA-GFP**, **CPCA-GFP** and **GPCA-GFP**. Mouse select image above left for larger view.



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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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Dr—D. rerio Dm—D. melanogaster Sm—S. mutans Ce—C. elegans Sc—S. cerevisiae Sa—S. aureus Ec—E. coli.*