abcam

Product datasheet

Recombinant A. victoria GFP protein ab84191

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Description

Product name Recombinant A. victoria GFP protein

Purity > 95 % SDS-PAGE.

Purified using conventional chromatography techniques.

Endotoxin level < 1.000 Eu/μg
Expression system Escherichia coli

Accession P42212

Protein length Full length protein

Animal free No

Nature Recombinant

Species Aequorea victoria

Sequence MSKGEELFTG VVPILVELDG DVNGHKFSVS

GEGEGDATYG KLTLKFICTT GKLPVPWPTL VTTFSYGVQC FSRYPDHMKQ HDFFKSAMPE

GYVQERTIFF KDDGNYKTRA EVKFEGDTLV NRIELKGIDF KEDGNILGHK LEYNYNSHNV YIMADKQKNG IKVNFKIRHN

IEDGSVQLAD HYQQNTPIGD GPVLLPDNHY

LSTQSALSKD PNEKRDHMVL LEFVTAAGIT HGMDELYK

Predicted molecular weight 27 kDa

Amino acids 1 to 238

Additional sequence information (AAA27721) MW confirmed by MALDI-TOF.

Description Recombinant A. victoria GFP protein

Specifications

Our Abpromise guarantee covers the use of ab84191 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications SDS-PAGE

Western blot

Functional Studies

Mass spectrometry MALDI-TOF

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Form Liquid

Additional notes Endotoxin Level determined by LAL method.

Concentration determined by Bradford assay.

Preparation and Storage

Stability and Storage

Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -80°C. Avoid freeze / thaw cycle.

pH: 8.00

Constituents: 0.316% Tris HCI, 10% Glycerol

This product is an active protein and may elicit a biological response in vivo, handle with caution.

General Info

Relevance

Function: Energy-transfer acceptor. Its role is to transduce the blue chemiluminescence of the protein aequorin into green fluorescent light by energy transfer. Fluoresces in vivo upon receiving energy from the Ca²⁺ -activated photoprotein aequorin.

Subunit structure: Monomer.

Tissue specificity: Photocytes.

Post-translational modification: Contains a chromophore consisting of modified amino acid residues. The chromophore is formed by autocatalytic backbone condensation between Ser-65 and Gly-67, and oxidation of Tyr-66 to didehydrotyrosine. Maturation of the chromophore requires nothing other than molecular oxygen.

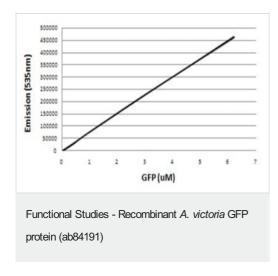
Biotechnological use: Green fluorescent protein has been engineered to produce a vast number of variously colored mutants, fusion proteins, and biosensors. Fluorescent proteins and its mutated allelic forms, blue, cyan and yellow have become a useful and ubiquitous tool for making chimeric proteins, where they function as a fluorescent protein tag. Typically they tolerate N- and C-terminal fusion to a broad variety of proteins. They have been expressed in most known cell types and are used as a noninvasive fluorescent marker in living cells and organisms. They enable a wide range of applications where they have functioned as a cell lineage tracer, reporter of gene expression, or as a measure of protein-protein interactions. Can also be used as a molecular thermometer, allowing accurate temperature measurements in fluids. The measurement process relies on the detection of the blinking of GFP using fluorescence correlation spectroscopy.

Sequence similarities: Belongs to the GFP family.

Biophysicochemical properties: Absorption: Abs(max)=395 nm

Exhibits a smaller absorbance peak at 470 nm. The fluorescence emission spectrum peaks at 509 nm with a shoulder at 540 nm.

Images



- 1. Prepare a 150 μ I recombinant GFP with various concentrations (0.00074 nM 6.21 μ M) in assay buffer and equilibrate to 25°C. (Assay buffer: 10 mM Tris-HCI (pH 8.0), 10 mM EDTA, 0.02% sodium azide.)
- 2. Read at excitation wavelengths 485 nm and emission 535 nm.
- 96 Well Polystyrene Microplate, black
- Fluorescent plate reader



3 ug of reduced ab84191 on SDS-PAGE, stained with Coomassie Blue.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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