



Product Information

Aequorin from jellyfish (*Aequorea* sp.)

Product Number **A 4140**

Storage Temperature -0 °C

Product Description

CAS Number: 50934-79-7

Molecular Weight: 21.7 kDa

pI: 4.2 - 4.9

Aequorin is one of a group of photoproteins isolated from marine coelenterates which emit blue light in the presence of Ca^{2+} ions. This bioluminescence is somewhat unique in that molecular oxygen is not required for light emission. The chromophore, coelenterazine,¹ is bound to aequorin and the binding of calcium triggers the oxidation of the chromophore, producing a photon of light with a wavelength of 470 nm. *In vivo*, many organisms have an associated green fluorescent protein which shifts the emission to higher wavelengths, making the color of the emitted light appear green.²

Aequorin from *Aequorea aequorea* has been purified to homogeneity and has been sequenced from cDNA cloning into various hosts. It is an 189 amino acid protein formed into a hydrophobic core region where coelenterazine is bound as what is believed to be a peroxidized form, since molecular oxygen is not required for light emission. Crystallography studies of the protein have elucidated 3 possible binding sites for Ca^{2+} , but only 1 of these is absolutely required for light output.³

Because of the high sensitivity which can be achieved in photon detection systems, aequorin has been extensively studied as a means of quantitation of cellular calcium in various biological systems. The advantages of such a system are high sensitivity, relative sensitivity for Ca^{2+} , ease of signal detection, and lack of toxicity in biological systems. However, these desirable properties are offset by several difficulties encountered in experimental design and data collection: scarcity of purified proteins, large molecular size, one-time reactivity, influence of experimental conditions on sensitivity, nonlinearity of the relation between Ca^{2+} concentration and light intensity, and limited speed of response in light intensity to changes in Ca^{2+} concentration.

In spite of these difficulties, aequorin has been shown to be an effective tool in analysis of Ca^{2+} concentrations and flux in biological systems. Measurements of Ca^{2+} in *Xenopus*,⁴ *E. coli*,⁵ and mammalian cells⁶ have demonstrated the utility of photometric measurement of Ca^{2+} . Molecular biology techniques have been used to develop aequorin fusion proteins for intracellular measurements and localization of Ca^{2+} in plasma membranes⁷ and cytoplasm,^{8,9} and to develop detection systems for proteolytic activity in cells.¹⁰

Precautions and Disclaimer

For Laboratory Use Only. Not for drug, household or other uses.

Preparation Instructions

This product is soluble in water (10 mg/ml), yielding a clear colorless solution.

References

1. Shimomura, O., and Johnson, F., Properties of the bioluminescent protein aequorin. *Biochemistry*, **8(10)**, 3991-3997 (1969).
2. Blinks, J. R., et al., Photoproteins as biological calcium indicators. *Pharmacological Reviews*, **28(1)**, 1-93 (1976).
3. Head, J. F., et al., The crystal structure of the photoprotein aequorin at 2.3 Å resolution. *Nature*, **405(6784)**, 372-376 (2000).
4. Grygorczyk, R., et al., Detection of intracellular calcium elevations in *Xenopus laevis* oocytes: aequorin luminescence versus electrophysiology. *J. Neurosci. Methods*, **67(1)**, 19-25 (1996).
5. Jones, H. E., et al., Direct measurement of free Ca^{2+} shows different regulation of Ca^{2+} between the periplasm and the cytosol of *Escherichia coli*. *Cell Calcium*, **32(4)**, 183-192 (2002).
6. Brandenburger, Y., et al., Measurement of perimitochondrial Ca^{2+} concentration in bovine adrenal glomerulosa cells with aequorin targeted to the outer mitochondrial membrane. *Biochem. J.*, **341(Pt 3)**, 745-753 (1999).

7. Nakahashi, Y., et al., Construction of a full-length Ca^{2+} -sensitive adenylyl cyclase/aequorin chimera. J. Biol. Chem., **272(29)**, 18093-18097 (1997).
8. Shaw, B. D., et al., Expression of recombinant aequorin as an intracellular calcium reporter in the phytopathogenic fungus *Phylosticta ampellicida*. Fungal Genet. Biol., **34(3)**, 207-215 (2001).
9. Chandra, S., et al., Measurement of Ca^{2+} fluxes during elicitation of the oxidative burst in aequorin-transformed tobacco cells. J. Biol. Chem., **272(45)**, 28274-28280 (1997).
10. Waud, J. P., et al., Measurement of proteases using chemiluminescence-resonance-energy-transfer chimeras between green fluorescent protein and aequorin. Biochem. J., **357(Pt 3)**, 687-697 (2001).

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