# Rubisco activase (aA-18): sc-15864



The Power to Question

### **BACKGROUND**

The Calvin cycle uses ATP and NADPH generated by the light reactions of photosynthesis to reduce carbon dioxide to sugar. The enzyme that catalyzes the first step in the Calvin cycle is ribulose bis-phosphate carboxylase (designated Rubisco). Rubisco activase is a nuclear encoded protein that is transported to the chloroplasts where it catalyzes the activation of Rubisco. The *Arabidopsis* Rubisco activase gene, which is located on chromosome 2, encodes mRNA that undergoes alternative splicing to yield two forms of Rubisco activase protein. Rubisco activase is expressed throughout the green parts of the plant, but not in roots or petals. Rubisco activase protein content accounts for approximately 5% of the total soluble protein in the green plant tissue. Expression of Rubisco activase is regulated by light, carbon-dioxide and a circadian clock.

# **REFERENCES**

- Werneke, J.M., Zielinski, R.E. and Ogren, W.L. 1988. Structure and expression of spinach leaf cDNA encoding ribulosebisphosphate carboxylase/oxygenase activase. Proc. Natl. Acad. Sci. USA 85: 787-791.
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- 6. Cheng, S.H., Moore, B. and Seemann, J.R. 1998. Effects of short and long-term elevated  $\mathrm{CO}_2$  on the expression of ribulose-1, 5-bisphosphate carboxylase/oxygenase genes and carbohydrate accumulation in leaves of *Arabidopsis thaliana* (L.) Heynh. Plant Physiol. 116: 715-723.
- Lin, X., Kaul, S., Rounsley, S., Shea, T.P., Benito, M.I., Town, C.D., Fujii, C.Y., Mason, T., Bowman, C.L., Barnstead, M., Feldblyum, T.V., Buell, C.R., Ketchum, K.A., Lee, J., Ronning, C.M., Koo, H.L., Moffat, K.S., Cronin, L.A., et al. 1999. Sequence and analysis of chromosome 2 of the plant *Arabidopsis thaliana*. Nature 402: 761-768.

# **SOURCE**

Rubisco activase (aA-18) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the C-terminus of Rubisco activase of *Arabidopsis thaliana* origin.

#### **STORAGE**

Store at  $4^{\circ}$  C, \*\*DO NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

#### **PRODUCT**

Each vial contains 200  $\mu g$  IgG in 1.0 ml of PBS with <0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-15864 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

### **APPLICATIONS**

Rubisco activase (aA-18) is recommended for detection of Rubisco activase of *Arabidopsis thaliana, Lycopersicon esculentum* and *Nicotiana tabacum* origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

#### **RECOMMENDED SECONDARY REAGENTS**

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluorescence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

# **SELECT PRODUCT CITATIONS**

- Chen, G., Bi, Y.R. and Li, N. 2005. EGY1 encodes a membrane-associated and ATP-independent metalloprotease that is required for chloroplast development. Plant J. 41: 364-375.
- 2. Yin, Z., Meng, F., Song, H., Wang, X., Xu, X. and Yu, D. 2009. Expression quantitative trait loci analysis of two genes encoding Rubisco activase in soybean. Plant Physiol. 152: 1625-1637.

## **RESEARCH USE**

For research use only, not for use in diagnostic procedures.

### **PROTOCOLS**

See our web site at www.scbt.com or our catalog for detailed protocols and support products.

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