

This product is for research use only (not for diagnostic or therapeutic use)

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## Product no AS10 690-DL594

# Anti-Clathrin heavy-chain 1,2, DyLight® 594 conjugated (40 μg)

#### **Product information**

Immunogen

KLH-conjugated peptide derived from available plant clathrin heavy chain sequences including Arabidopsis thaliana clathrin heavy chain 1 UniProt: Q0WNJ6, TAIR:At3q11130, clathrin heavy chain 2 UniProt: Q0WLB5,

**Host** Rabbit

Clonality Polyclonal

**Purity** Immunogen affinity purified serum, in PBS pH 7.4, conjugated to DyLight® 594.

Format Liquid in PBS pH 7,4.

Quantity 40 μg

Storage Store at 4°C for 12-18 months, A preservative may be added for long time storage up to 2 years. Spin briefly the tube

before use

Additional information DyLight® 594 has Amax = 593 nm, Emax = 618 nm. DyLight® is a registered trademark of Thermofisher Inc., and its

subsidiaries.

### **Application information**

**Recommended dilution** To be determined by end user.

Expected | apparent MW 193 | 170 kDa (Arabidopsis thaliana)

Confirmed reactivity Zea mays

Predicted reactivity Amborella trichopoda, Arabidopsis thaliana, Brassica napus, Capsella rubella, Chlamydomonas reinhardtii, Citrus

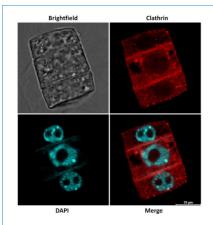
aurantium var. sinensis, Eucalyptus grandis, Glycine max, Chlorella variabilis, Leucaena glauca, Lotus japonicus, Medicago tribuloides, Mimulus guttatus, Musa malaccensis, Nicotiana tabacum, Oryza sativa, Panicum italicum, Physcomitrium patens, Phaseolus vulgaris, Pisum sativum, Populus balsamifera, Populus trichocarpa, Ricinus communis, Selaginella moellendorffii, Sisymbrium salsugineum, Solanum lycopersicum, Theobroma cacao, Triticum

aestivum, Vitis vinifera, Zea mays.

Species of your interest not listed? Contact us

Not reactive in Nicotiana benthamiana

**Selected references** To be added when available. Antibody released in May 2023.



Material: Zea mays hybrid variety

#### Preparation

5 days-old germinating maize root tips were cut and fixed with 4% formaldehyde, 60 min, RT. Cell walls were digested with enzymes cellulase and pectinase in MESbuffer for 90 min, RT. After washing with PBS, roots were squashed gently using a flat tip forceps to release the cells into PBS buffer. Cells were allowed to settle O/N at 4°C, followed by immunolocalization.

#### Immunocytochemistry (ICC)

Cells were permeabilized with 0.5% Triton X-100, 10 min, RT, followed by washing with PBS buffer before blocking with 5% fish gelatin-PBS, 30



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min, RT.

**Detection antibody**: Cells were incubated with rabbit anti-clathrin 1,2 primary antibodies conjugated with DyLight®594 (AS10 690-DL594, Agrisera) for 3h/RT in the dark at 1:100 dilution. Nuclei were stained with DAPI followed Fluoromount-G mounting (Southern Biotech).

Courtesy of Dr. Ferhan Ayaydin and Dr. Divya Teja Dondapati, Hungarian Centre of Excellence for Molecular Medicine, (HCEMM), Szeged, Hungary.