

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

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Product no AS15 3091

Anti-VDE | Violaxanthin de-epoxidase

Product information

Immunogen His-tagged, recombinant VDE from Arabidopsis thaliana, overexpressed in E.coli, Uniprot: Q39249, TAIR: AT1G08550

Host Rabbit

Clonality Polyclonal

Purity Serum

Format Lyophilized

Quantity 50 μl

Reconstitution For reconstitution add 200 μl of sterile water

Store lyophilized/reconstituted at -20°C; once reconstituted make aliquots to avoid repeated freeze-thaw cycles. Please

remember to spin the tubes briefly prior to opening them to avoid any losses that might occur from material adhering to

the cap or sides of the tube.

Application information

Recommended dilution 1:2000 (WB)

Expected | apparent

Propeptide 52 kDa, mature protein: 39.7 kDa, migrates at 40 kDa (Arabidopsis thaliana)

Predicted reactivity

Brassica napus, Camellia sinensis, Citrus limon, Cucumis sativus, Chrysanthemum morifolium, Citrus sinensis, Coffea arabica, Fragaria ananassa, Glycine max, Glycine soja, Gossypium arboreum, Lactuca sativa, Lycium barbarum, Morus notabilis, Nicotiana tabacum, Phyllostachys edulis, Picea abies, Prunus humilis, Spinacia oleracea, Solanum lycopersicum, Theobroma cacao, Triticum aestivum, Zea mays, Zingiber officinale, Vitis vinifera

Species of your interest not listed? Contact us

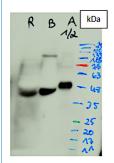
Not reactive in

Diatoms

Additional information

Extraction buffer must contain 7 M urea for Brachypodium distachyon and Oryza sativa samples.

This product can be sold containing ProClin if requested.



Samples:

R – 10 μL of total protein extract of *Oryza sativa* (50 mg macerated leaves in 100 μL Lysis Buffer).

B – 10 μL of total protein extract of Brachypodium distachyon (50 mg macerated leaves in 100 μL Lysis Buffer).

A – 10 μL of total protein extract of Arabidopsis thaliana diluted to 1:2 (50 mg macerated leaves in 200 μL 2X Laemmli Buffer). MW marker: NZYColour Protein Marker II from NZYTech.

10 μL of total protein extracted freshly from Arabidopsis thaliana, Brachypodium distachyon, and Oryza sativa with Lysis Buffer (7M Urea, 2M Thiourea, 4% CHAPS, 35 mM Tris), and denatured in 1X Laemmli with 25 mM DTT at 95°C for 4 min, were separated on 10 % SDS-PAGE and blotted 1h to PVDF membrane (pore size of 0.45 µm, GE Healthcare), using wet transfer. Blot was blocked with Blocking solution containing 5% milk in TBS-T, for 1h/RT with agitation. Blot was incubated in the primary antibody at a dilution of 1:2000 in Blocking solution containing 5% milk in TBS-T, for O/N at 4°C, with agitation. The antibody solution was decanted, and the blot was washed 3 times for 10 min in TBS-T at RT with agitation. Blot was incubated in secondary antibody (anti-rabbit IgG, horse radish peroxidase conjugated) diluted to 1:20 000 in Blocking solution containing 5% milk in TBS-T, for 1h/RT with agitation. The blot was washed 6 times for 5 min in TBS-T at RT with agitation and developed for 30 seconds with Agrisera ECLSuperBright. Exposure time was 10 seconds.

Courtesy of Dr. Nelson Saibo, Plant Gene Regulation Lab (GPlantS Unit), ITQB NOVA, Portugal