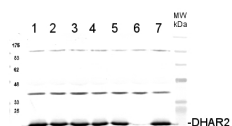


Product no **AS11 1747****DHAR2 | Dehydroascorbate Reductase 2****Product information**

<b>Immunogen</b>	KLH-conjugated synthetic peptide derived from known DHAR1 sequence of <i>Arabidopsis thaliana</i> <a href="#">Q9FRL8</a> , <a href="#">At1g75270</a>
<b>Host</b>	Rabbit
<b>Clonality</b>	Polyclonal
<b>Purity</b>	Immunogen affinity purified serum in PBS pH 7.4.
<b>Format</b>	Lyophilized
<b>Quantity</b>	200 µg
<b>Reconstitution</b>	For reconstitution add 200 µl of sterile water
<b>Storage</b>	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**

<b>Recommended dilution</b>	1 : 5000 (WB)
<b>Expected   apparent MW</b>	23,6   23,4 kDa
<b>Confirmed reactivity</b>	<i>Arabidopsis thaliana</i>
<b>Predicted reactivity</b>	<i>Ricinus communis</i> , <i>Populus trichocarpa</i> Species of your interest not listed? <a href="#">Contact us</a>
<b>Not reactive in</b>	No confirmed exceptions from predicted reactivity are currently known
<b>Selected references</b>	<a href="#">Grefen et al. (2009)</a> . The determination of protein-protein interactions by the mating-based split-ubiquitin system (mbSUS). <i>Methods Mol Biol</i> 479:217-233.

**application example**

1cm<sup>2</sup> of a leaf from *Arabidopsis thaliana* Col-0 (**1**) and or t-DNA insertion lines dhar1-1 (**2**), dhar1-2 (**3**), dhar1-3 (**4**), dhar2-1 (**5**), dhar2-2 (**6**), dhar1-3 EOS-DHAR1 (**7**), was extracted using 200µl Lyse&Load-Buffer (Grefen et al. 2009). 10 µl were separated on a 15% SDS-PAGE and blotted 1h to PVDF (using Bjerrum Buffer in a semidry blot). Blots were blocked with 5% Milk in 1xTBS-Tween20 (1%) for 1h at room temperature (RT) with agitation. Blot was incubated in the primary antibody at a dilution of 1:5000 (in 5% Milk 1xTBS-Tween20 (1%) + 0.01 % NaN<sub>3</sub>) ON at 4 °C with agitation. The antibody solution was decanted and the blot was washed 3 times for 10 minutes with 1x TBS-Tween20 at RT with agitation. Blot was incubated in secondary antibody BioRad anti-rabbit IgG AP-conjugate (#170-6518) diluted to 1:2000 in 5% Milk 1xTBS-Tween20 (1%) + 0.01 % NaN<sub>3</sub> for 1h at RT with agitation. The blot was washed as above, equilibrated in staining buffer (100mM Tris-HCl, 100mM NaCl, 5mM MgCl<sub>2</sub>, see Grefen et al. 2009) and developed for 5-15 min. with staining solution (Nitro blue tetrazolium chloride (NBT) and 5-bromo-4-chloro-3-indoylphosphate-p-toluidin (BCIP) in staining buffer).

Courtesy Dr. Chrisopher Grefen, UK