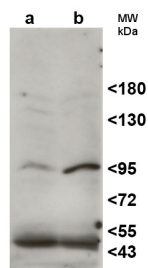


Product no **AS15 3095****HEN1 | HUA ENHANCER 1****Product information**

| | |
|-----------------------|---|
| Immunogen | KLH-conjugated peptide derived from <i>Arabidopsis thaliana</i> HEN1 sequence, Uniprot: Q9C5Q8 , TAIR: AT4G20910 |
| Host | Rabbit |
| Clonality | Polyclonal |
| Purity | Immunogen affinity purified serum in PBS pH 7.4. |
| Format | Lyophilized |
| Quantity | 50 µg |
| Reconstitution | For reconstitution add 50 µl of sterile water |
| Storage | 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 : 1000-1 : 5000 (WB) |
| Expected apparent MW | 104,5 kDa 105 kDa |
| Confirmed reactivity | <i>Arabidopsis thaliana</i> |
| Not reactive in | No confirmed exceptions from predicted reactivity are currently known |

application example

50 µg of total protein from *Arabidopsis thaliana* whole vegetative rosette wild type Col-0 (**a**) and HEN1 overexpression mutant (**b**) extracted with extraction buffer (50 mM Tris pH 7.5; 150 mM NaCl; 1 mM EDTA; 10 % v/v Glycerin; 1 mM DTT, 1x Complete Protease Inhibitor Cocktail, Roche) and denatured with Laemmli buffer at 95°C/5 min., were separated on 7.5 % SDS-PAGE and blotted 1.5 h to PVDF using tank transfer. Blots were blocked with blocking buffer (5% milk powder; 1x TBS; 0.1% Tween-20) overnight at 4°C with agitation. Blot was incubated in the primary antibody at a dilution of 1:1000 for 2h at RT with agitation. The antibody solution was decanted and the blot was rinsed briefly and then washed three times for 15 min in TBS-T at RT with agitation. Blot was incubated in secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated, from Agrisera, [AS09 602](#)) diluted to 1:20 000 in blocking buffer for 1h at RT with agitation. The blot was washed as above and developed for 5 min with Amersham ECL Prime and expose to Amersham Hyperfilms ECL for 20 seconds.

Courtesy of Dr. Pablo Manavella, Instituto de Agrobiotecnología del Litoral (IAL), Argentina