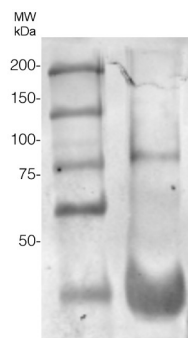


Product no **AS10 672****Anti-AGO6 | Argonaute 6****Product information****Immunogen** | KLH-conjugated peptide derived from *Arabidopsis thaliana* AGO6 UniProt: [Q48771](#) TAIR: [At2g32940](#)**Host** | Rabbit**Clonality** | Polyclonal**Purity** | Immunogen affinity purified serum in PBS pH 7.4.**Format** | 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.**Additional information** | There can be a reaction with NEB broad range prestained protein ladder depending upon secondary antibody used.AGO6 accumulates to lower levels in the *nprp1a* mutant ([Havecker](#) et al. 2010). In addition, a non-specific band migrates to a similar location and thus protein was run on a 6% denaturing gel to obtain good separation.

If other secondary antibody is used than in the figure below, cross-reacting pattern can be different.

Application information**Recommended dilution** | 1 : 1000 (WB)**Expected | apparent MW** | 116.4 | 99 kDa**Confirmed reactivity** | *Arabidopsis thaliana***Predicted reactivity** | *Arabidopsis thaliana***Not reactive in** | *Zea mays***Additional information** | AGO6 expression is cell-specific. Seedlings can be used as a negative control**Selected references** | [Oliver](#) & Martinez. (2021) Accumulation dynamics of ARGONAUTE proteins during meiosis in Arabidopsis. Plant Reprod. 2021 Nov 23. doi: 10.1007/s00497-021-00434-z. Epub ahead of print. PMID: 34812935.
[Sprunck](#) et al. (2019). Elucidating small RNA pathways in Arabidopsis thaliana egg cells. <http://dx.doi.org/10.1101/525956>
[Havecker](#) et al. (2010). The RNA-directed DNA methylation Arabidopsis Argonautes functionally diverge based on expression and interaction with target loci. Plant Cell. 2010 Feb;22(2):321-34. doi: 10.1105/tpc.109.072199.**Application example**

360 µg/well of *Arabidopsis thaliana* protein extracted from leaf using TCA-protein precipitation (for method check protocol tab) and saturated in 8M urea were separated on 15% SDS-PAGE and blotted for 1 hour to 0.2 µm nitrocellulose at 100V using wet transfer system. Blots were blocked with 0.5% cold fish gelatin for 1hr at room temp with agitation. Blot was incubated in the primary antibody (anti-H3) at a dilution of 1:2500 for an hour at RT with agitation. The blots were washed with 3X 15min TBS-TT at RT with agitation. Blots as incubated in the secondary antibody (goat anti-rabbit DyLight® 800 conjugated, [AS12 2460](#), Agrisera) 1:5000 dilution for 30min at RT with agitation and washed 1X with TBSTT for 15min,



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1X with TBST for 15min before scanning with the ODyssey IRD scanner.

Courtesy of Dr. Betty Chung and Pawel Baster, University of Cambridge, United Kingdom