

# Agrisera

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

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Product no **AS12 2634**

## DRP5B | Dynamin related protein 5B

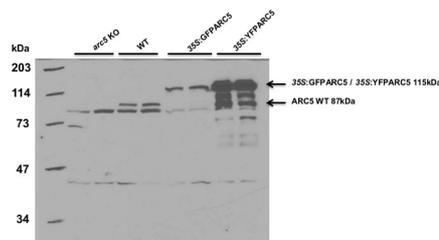
### Product information

<b>Immunogen</b>	KLH-conjugated synthetic peptide derived from <i>Arabidopsis thaliana</i> DRP5B sequence UniProt: <a href="#">Q84N64</a> , TAIR: <a href="#">AT3G19720</a>
<b>Host</b>	Rabbit
<b>Clonality</b>	Polyclonal
<b>Purity</b>	Affinity purified serum in PBS, pH 7.4
<b>Format</b>	Lyophilized
<b>Quantity</b>	50 µg
<b>Reconstitution</b>	For reconstitution add 50 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 tubes briefly prior to opening them to avoid any losses that might occur from lyophilized material adhering to the cap or sides of the tubes.

### Application information

<b>Recommended dilution</b>	1 : 5000 (WB)
<b>Expected   apparent MW</b>	87.17 kDa
<b>Confirmed reactivity</b>	<i>Arabidopsis thaliana</i>
<b>Predicted reactivity</b>	<i>Glycne max</i> , <i>Manihot esculenta</i> , <i>Mesostigma viride</i> , <i>Oryza sativa</i> , <i>Physcomitrella patens</i> , <i>Sorghum bicolor</i> , <i>Triticum aestivum</i> , <i>Zea mays</i> , <i>Vitis vinifera</i> Species of your interest not listed? <a href="#">Contact us</a>
<b>Not reactive in</b>	<i>Chlamydomonas reinhardtii</i>
<b>Additional information</b>	Chemiluminescent <a href="#">extreme low femtogram detection</a> range reagent is recommended to use with this antibody. For high resolution images, please visit the specific product page at <a href="http://www.agrisera.com">www.agrisera.com</a>

#### Application example



Total proteins extracted from 5 mg leaf or flower bud of *Arabidopsis thaliana* tissue with SDS-loading buffer were separated on 10% SDS-PAGE using tank transfer and blotted for 1 h to nitrocellulose membrane. Blots were blocked with 2% non-fat dry milk for 1 h at room temperature (RT) with agitation. Blot was incubated in the primary antibody (affinity-purified DRP5B) at a dilution of 1:5 000 in TBS-T + 2% non-fat dry milk overnight at RT with agitation. The antibody solution was decanted and the blot was rinsed briefly, then washed 4 times for 10 min in TBS-T at RT with agitation. Blot was incubated in secondary antibody (anti-rabbit IgG horseradish peroxidase conjugated, from Agrisera [AS09 602](#)) diluted to 1:5 000 in TBS-T + 2% non-fat dry milk for 2 hr at RT with agitation. The blot was washed as above and developed for 5 min with SuperSignal West Pico (Thermo Scientific) according to the manufacturer's instructions. Exposure time was 30 min. Additionally the same blot was washed and developed with extreme low femtogram detection range chemiluminescent reagent. Exposure time was 1 min. The latter helped to increase DRP5B detection with the affinity-purified DRP5B antibodies. The signal was not obtained from bud tissue, suggesting that expression level of

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DRP5B may be too low for detection (blot not included).

Courtesy of Deena Kadirjan-Kalbach and Katherine W. Osteryoung, Michigan State University, USA