



## PIN1 (aK-13): sc-27161

### BACKGROUND

Polar auxin transport controls multiple developmental processes in plants, including the formation of vascular tissue. Mutations affecting the PIN-FORMED (PIN1) gene diminish polar auxin transport in *Arabidopsis thaliana* inflorescence axes. The PIN1 gene encodes a 67 kDa protein with similarity to bacterial and eukaryotic carrier proteins. The PIN1 protein is at the basal end of auxin transport-competent cells in vascular tissue and may act as a transmembrane component of the auxin efflux carrier.

### REFERENCES

- Galweiler, L., Guan, C., Muller, A., Wisman, E., Mendgen, K., Yephremov, A., Palme, K. 1998. Regulation of polar auxin transport by AtPIN1 in *Arabidopsis* vascular tissue. *Science* 282: 2226-2230.
- Steinmann, T., Geldner, N., Grebe, M., Mangold, S., Jackson, C.L., Paris, S., Galweiler, L., Palme, K., Jurgens, G. 1999. Coordinated polar localization of auxin efflux carrier PIN1 by GNOM ARF GEF. *Science*. 286: 316-8.
- Swarup, R., Marchant, A., Bennett, M.J., 2000. Auxin transport: providing a sense of direction during plant development. *Biochemical Soc. Trans.* 28: 481-485.
- Geldner, N., Friml, J., Stierhof, Y.D., Jurgens, G., Palme, K. 2001. Auxin transport inhibitors block PIN1 cycling and vesicle trafficking. *Nature*. 413: 425-8.
- Noh, B., Bandyopadhyay, A., Peer, W.A., Spalding, E.P., Murphy, A.S., 2003. Enhanced gravi- and phototropism in plant mdr mutants mislocalizing the auxin efflux protein PIN1. *Nature*. 423: 999-1002.
- Uchida, T., Takamiya, M., Takahashi, M., Miyashita, H., Ikeda, H., Terada, T., Matsuo, Y., Shirouzu, M., Yokoyama, S., Fujimori, F., Hunter, T., 2003. Pin1 and Par14 peptidyl prolyl isomerase inhibitors block cell proliferation. *Chem. Biol.* 10: 15-24.
- Blakeslee, J.J., Bandyopadhyay, A., Peer, W.A., Makam, S.N., Murphy, A.S., 2004. Relocalization of the PIN1 auxin efflux facilitator plays a role in phototropic responses. *Plant Physiol.* 134: 28-31.

### SOURCE

PIN1 (aK-13) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the N-terminus of PIN1 of *Arabidopsis thaliana* origin.

### PRODUCT

Each vial contains 200 µg IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-27161 P, (100 µg peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

### STORAGE

Store at 4° C, \*\*DO NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

### RESEARCH USE

For research use only, not for use in diagnostic procedures.

### APPLICATIONS

PIN1 (aK-13) is recommended for detection of PIN1 of *Arabidopsis thaliana* origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Molecular Weight of PIN1: 67 kDa.

### RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluorescence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

### PROTOCOLS

See our web site at [www.scbt.com](http://www.scbt.com) or our catalog for detailed protocols and support products.