Phospho-PAR-4 (Thr163) Antibody



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Applications: WB	Reactivity:	Sensitivity: Endogenous	MW (kDa): 43	Source: Rabbit	UniProt ID: #Q96IZ0	Entrez-Gene Id 5074
Product Usage Information	Application			Dilution		
	We	Western Blotting			1:1000	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity / Sensit			,	endogenous levels of PAR-4 when phosphorylated at Thr163 and is equivalent to Thr155 in rat and Thr156 in mouse).		
Species predicted react based on 100 sequence homology	0%	Mouse, Rat, Monkey				

Source / Purification

Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Thr163 of human PAR-4 (Thr155 in rat and Thr156 in mouse). Antibodies were purified by protein A and peptide affinity chromatography.

Background

PAR-4 (prostate apoptosis response-4) was identified as a protein that is upregulated in prostate tumor cells undergoing apoptosis (1). Additionally, in parallel studies PAR-4 was found in the yeast two-hybrid system to bind to the Wilms' tumor suppressor protein WT1 and may modulate WT1-medated transcriptional activation (2). PAR-4 contains a leucine zipper domain and a death domain and has been implicated as an effector of apoptosis during tumorigenesis as well as in neurodegenerative disorders (3,4). PAR-4 is widely expressed in normal tissues but can be downregulated in some tumor types. The mechanism of PAR-4 mediated apoptosis regulation appears to be complex and dependent on the cellular context. Studies have indicated roles for PAR-4 in activation of the Fas-FADD-caspase-8 pathway as well as inhibition of the NF-kB pro-survival pathway (5-7). Its activity is likely to depend on the cellular context and post-translational modifications. For instance, phosphorylation of PAR-4 by Akt prevents its nuclear translocation thereby promoting cell survival (8). In contrast, phoshorylation of rat PAR-4 at T155 by PKA appears to positively regulate its apoptotic activity (9).

Background References

- 1. Sells, S.F. et al. (1997) Mol. Cell Biol. 17, 3823-3832.
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- 3. Guo, Q. et al. (1998) Nat. Med. 4, 957-962.
- 4. El-Guendy, N. and Rangnekar, V.M. (2003) Exp. Cell Res. 283, 51-66.
- 5. Chakraborty, M. et al. (2001) Cancer Res. 61, 7255-7263.
- 6. Díaz-Meco, M.T. et al. (1996) Cell 86, 777-786.
- 7. Diaz-Meco, M.T. et al. (1999) *J. Biol. Chem.* 274, 19606-79612.
- 8. Goswami, A. et al. (2005) Mol. Cell 20, 33-44.
- 9. Gurumurthy, S. et al. (2005) Mol. Cell Biol. 25, 1146-1161.

Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key

Cross-Reactivity Key

WB: Western Blotting

Phospho-PAR-4 (Thr163) Antibody (#2329) Datasheet Without Images Cell Signaling Technology

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse GP: Guinea Pig Rab: rabbit All: all species expected

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