PAR-4 Ab

Cat.#: AF4785 Concn.: 1mg/ml Mol.Wt.: 43kDa Size: 50ul,100ul,200ul Source: Rabbit Clonality: Polyclonal

Application: WB 1:500-1:2000

Reactivity: Human, Mouse, Rat

Purification: The antiserum was purified by peptide affinity

chromatography using SulfoLink™ Coupling Resin (Thermo

Fisher Scientific).

Specificity: PAR-4 Ab detects endogenous levels of PAR-4.

Immunogen: A synthesized peptide derived from human PAR-4.

Uniprot: Q96IZ0

Subcellular Location: Cytoplasm. Nucleus. Mainly cytoplasmic in absence of

apoptosis signal and in normal cells. Nuclear in most cancer cell lines. Nuclear entry seems to be essential but not sufficient for apoptosis (By similarity). Nuclear localization

includes nucleoplasm and PML nuclear bodies.

Tissue Specificity: Widely expressed. Expression is elevated in various

neurodegenerative diseases such as amyotrophic lateral sclerosis, Alzheimer, Parkinson and Huntington diseases and

stroke. Down-regulated in several cancers.

Similarity: The leucine-zipper domain is not essential for apoptosis, but

is required for sensitization of cells to exogenous apoptotic insults and for interaction with its partners. The SAC domain is a death-inducing domain selective for apoptosis induction in cancer cells. This domain is essential for nuclear entry, Fas activation, inhibition of NF-kappa-B activity and induction of apoptosis in cancer cells (By similarity). The B30.2/SPRY domain-binding motif mediates recognition by

proteins containing a B30.2/SPRY domain.

Storage Condition and

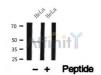
Buffer:

Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at -20

°C.Stable for 12 months from date of receipt.



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Western blot analysis of PAR-4 in lysates of HeLa , using PAR-4 Ab(AF4785).

<code>IMPORTANT:</code> For western blot, incubate membrane with diluted primary Ab in 5% w/v milk , 1X TBS, 0.1% Tween®20 at 4°C with gentle shaking, overnight.

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