

Product datasheet for AM00104PU-N

OriGene Technologies, Inc.

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PIN1 (incl. pos. control) Mouse Monoclonal Antibody [Clone ID: 8C10]

Product data:

Product Type: Primary Antibodies

Clone Name: 8C10

Applications: ELISA, WB

Recommended Dilution: ELISA: 0.1 µg/ml (protein ELISA).

Western Blot: 0.5 μg/ml for HRPO/ECL detection.

Recommended blocking buffer: Casein/Tween 20 based blocking and blot incubation buffer.

Included Positive Control: Cell lysate from untreated HepG2 cells.

Reactivity: Canine, Human, Mouse

Host: Mouse Isotype: IgG1

Clonality: Monoclonal

Immunogen: Recombinant Human PIN1

Specificity: This antibody specifically recognizes Human PIN1 in cell extracts at 16 kDa.

Formulation: 1 ml 2 x PBS containing 0.09% Sodium Azide, PEG and Sucrose

State: Purified

State: Lyophilized purified IgG fraction

Reconstitution Method: Restore with 1 ml H₂O (15 min, RT).

Purification: Purified from serum-free cell culture supernatant by Subsequent Thiophilic Adsorption and

Size Exclusion Chromatography

Conjugation: Unconjugated

Storage: Store lyophilized (preferably in a desiccator) at -20°C and reconstituted (aliquote and freeze

in liquid nitrogen) at -80°C.

Avoid repeated freezing and thawing.

Thaw aliquots at 37°C. Thawed aliquots may be stored at 4°C up to 3 months.

Stability: Shelf life: one year from despatch.

Gene Name: peptidylprolyl cis/trans isomerase, NIMA-interacting 1

Database Link: Entrez Gene 5300 Human

Q13526



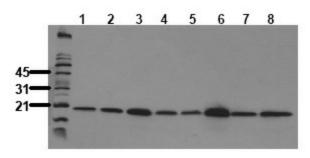


Background:

PIN1 is a peptidyl-prolyl-cis-trans-isomerase (PPlase) that specifically interacts with serine phosphate-proline or threonine phosphate-proline motifs. Upon binding, PIN1 isomerizes the peptide bond form cis to trans. Known substrates of PIN1 are serveral mitotic phosphoproteins (e.g. cdc25) as well as phosphorylated p53, phosphorylated β -catenin and phosphorylated tau protein. It is assumed that the isomerization of phosphoproteins regulates their biological function.

Synonyms: PIN-1

Product images:



Detection of endogenous PIN1 Whole cell lysates of serum starved tumor cells (20.000 cells per lane) were applied to SDS-PAGE and transferred to PVDF membranes. Immunoblots were probed with mab PIN1 8C10 (0.5 ug/ ml) for 1h at RT and developed by ECL (exp. time: 30 sec). lane 1: A431; lane 2: A549; lane 3: SKOV3; lane 4: OVCAR5; lane 5: HaCaT; lane 6: PC3; lane 7: HeLa; lane 8: HepG2