Store at -20C

Phospho-p57 Kip2 (Thr310) Antibody



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Applications: Reactivity: Sensitivity: MW (kDa): Source: **UniProt ID:** Entrez-Gene Id: WB, IF-IC Н Endogenous 57 Rabbit #P49918 1028

Product Usage Dilution Application Information 1:1000 Western Blotting 1:100 Immunofluorescence (Immunocytochemistry)

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% glycerol. Store at -**Storage** 20°C. Do not aliquot the antibody.

Phospho-p57 Kip2 (Thr310) Antibody detects endogenous levels of p57 Kip2 only when phosphorylated at

Specificity / Sensitivity threonine 310. This antibody may cross-react with p27 Kip1 when phosphorylated at Thr187.

Source / Purification Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Thr310 of human p57 Kip2. Antibodies are purified by protein A and peptide affinity

chromatography.

p27 Kip1 is a member of the Cip/Kip family of cyclin-dependent kinase inhibitors. Like its relatives, p57 **Background**

Kip2 and p21 Waf1/Cip1, the ability to enforce the G1 restriction point is derived from its inhibitory binding to CDK2/cyclin E and other CDK/cyclin complexes. Expression levels of p27 are upregulated in quiescent cells and in cells treated with cAMP or other negative cell cycle regulators. Downregulation of p27 can be induced by treatment with interleukin-2 or other mitogens; this involves phosphorylation of p27 and its

degradation by the ubiquitin-proteasome pathway (1-4).

Levels of p57 Kip2 are controlled by ubiquitination/degradation via the Skp1/Cul1/F-box-type E3 ubiquitin ligase complex SCF-Skp2, and this effect is dependent on Thr310 (5). A similar threonine phosphorylation

site in p27 Kip1, Thr187, has also been shown to regulate protein stability (6).

1. Lloyd, R.V. et al. (1999) Am J Pathol 154, 313-23. **Background References**

2. Polyak, K. et al. (1994) Genes Dev 8, 9-22.

3. Kato, J.Y. et al. (1994) Cell 79, 487-96.

4. Vlach, J. et al. (1997) EMBO J 16, 5334-44.

5. Kwon, T. et al. (2000) J Biol Chem 275, 423-8.

6. Heeneman, S. et al. (2000) J Biol Chem 275, 15926-32.

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

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, Western Blot Buffer

0.1% Tween® 20 at 4°C with gentle shaking, overnight.

WB: Western Blotting IF-IC: Immunofluorescence (Immunocytochemistry) **Applications Key**

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster **Cross-Reactivity Key**

X: Xenopus Z: zebrafish B: bovine Dq: dog Pq: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

GP: Guinea Pig Rab: rabbit All: all species expected

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