

**Phospho-mouse p21Cip1(S78) Antibody**  
**Affinity Purified Rabbit Polyclonal Antibody (Pab)**  
**Catalog # AP3877a****Specification**

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**Phospho-mouse p21Cip1(S78) Antibody - Product Information**

Application	DB,E
Primary Accession	<a href="#">P39689</a>
Other Accession	<a href="#">NP_001129489.1</a>
Reactivity	Mouse
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit IgG
Calculated MW	17785

**Phospho-mouse p21Cip1(S78) Antibody - Additional Information****Gene ID** 12575**Other Names**

Cyclin-dependent kinase inhibitor 1, CDK-interacting protein 1, Melanoma differentiation-associated protein, p21, Cdkn1a, Cip1, Waf1

**Target/Specificity**

This mouse p21Cip1 Antibody is generated from rabbits immunized with a KLH conjugated synthetic phosphopeptide corresponding to amino acid residues surrounding S78 of mouse p21Cip1.

**Dilution**

DB~~1:500

E~~Use at an assay dependent concentration.

**Format**

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

**Storage**

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions**

Phospho-mouse p21Cip1(S78) Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

**Phospho-mouse p21Cip1(S78) Antibody - Protein Information****Name** Cdkn1a

**Synonyms** Cip1, Waf1

**Function** May be involved in p53/TP53 mediated inhibition of cellular proliferation in response to DNA damage. Binds to and inhibits cyclin- dependent kinase activity, preventing phosphorylation of critical cyclin-dependent kinase substrates and blocking cell cycle progression. Functions in the nuclear localization and assembly of cyclin D-CDK4 complex and promotes its kinase activity towards RB1. At higher stoichiometric ratios, inhibits the kinase activity of the cyclin D- CDK4 complex (PubMed:[25329316](#)). Inhibits DNA synthesis by DNA polymerase delta by competing with POLD3 for PCNA binding (By similarity). Plays an important role in controlling cell cycle progression and DNA damage-induced G2 arrest (By similarity).

**Cellular Location**

Cytoplasm. Nucleus

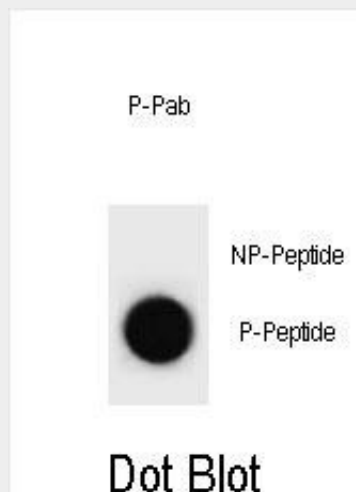
**Tissue Location**

Expressed in keratinocytes (at protein level).

**Phospho-mouse p21Cip1(S78) Antibody - Protocols**

Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

**Phospho-mouse p21Cip1(S78) Antibody - Images**

Dot blot analysis of Mouse p21Cip1 Antibody (Phospho S78) Phospho-specific Pab (Cat. #AP3877a) on nitrocellulose membrane. 50ng of Phospho-peptide or Non Phospho-peptide per dot were adsorbed. Antibody working concentrations are 0.6ug per ml.

**Phospho-mouse p21Cip1(S78) Antibody - Background**

The protein encoded by this gene belongs to the highly

conserved cyclin family, whose members are characterized by a dramatic periodicity in protein abundance through the cell cycle. Cyclins function as regulators of CDK kinases. Different cyclins exhibit distinct expression and degradation patterns which contribute to the temporal coordination of each mitotic event. This cyclin forms a complex with and functions as a regulatory subunit of CDK4 or CDK6, whose activity is required for cell cycle G1/S transition. This protein has been shown to interact with and be involved in the phosphorylation of tumor suppressor protein Rb. The CDK4 activity associated with this cyclin was reported to be necessary for cell cycle progression through G2 phase into mitosis after UV radiation. Several transcript variants encoding different isoforms have been found for this gene.

#### **Phospho-mouse p21Cip1(S78) Antibody - References**

Liu, C.Y., et al. Carcinogenesis 31(7):1259-1263(2010)  
Kim, J., et al. Cytokine 50(1):42-49(2010)  
Kamatani, Y., et al. Nat. Genet. 42(3):210-215(2010)  
Gumina, M.R., et al. Cell Cycle 9(4):820-828(2010)  
Radulovich, N., et al. Mol. Cancer 9, 24 (2010) :