

# Anti-MKI67IP (pT234) Antibody

Rabbit polyclonal antibody to MKI67IP (pT234) Catalog # AP60813

#### **Product Information**

Application WB, IF/IC, IHC
Primary Accession Q9BYG3
Other Accession Q91VE6

**Reactivity** Human, Mouse, Rat, Bovine

HostRabbitClonalityPolyclonalCalculated MW34222

#### **Additional Information**

**Gene ID** 84365

Other Names MKI67IP; NOPP34; MKI67 FHA domain-interacting nucleolar phosphoprotein;

Nucleolar phosphoprotein Nopp34; Nucleolar protein interacting with the

FHA domain of pKI-67; hNIFK

**Target/Specificity** Recognizes endogenous levels of MKI67IP (pT234) protein.

**Dilution** WB~~WB (1/500 - 1/2000), IHC (1/50 - 1/200), IF/IC (1/50 - 1/100) IF/IC~~N/A

IHC~~WB (1/500 - 1/2000), IHC (1/50 - 1/200), IF/IC (1/50 - 1/100)

**Format** Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30%

glycerol, and 0.09% (W/V) sodium azide.

**Storage** Store at -20 °C.Stable for 12 months from date of receipt

#### **Protein Information**

Name NIFK

Synonyms MKI67IP, NOPP34

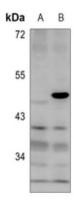
**Cellular Location** Nucleus, nucleolus. Chromosome. Note=Localizes to mitotic chromosomes in

conjunction with MKI67

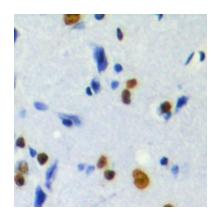
## **Background**

KLH-conjugated synthetic peptide encompassing a sequence within the center region of human MKI67IP. The exact sequence is proprietary.

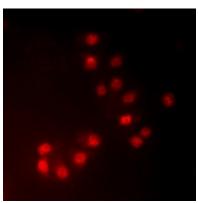
### **Images**



Western blot analysis of MKI67IP (pT234) expression in HGC27 (A), H446 (B) whole cell lysates.



Immunohistochemical analysis of MKI67IP (pT234) staining in human brain formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of MKI67IP (pT234) staining in HeLa cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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