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p16 INK4A (PT0242R) PT® Rabbit mAb

Isotype

IgG,Kappa

CatalogNo: YM8152 Recombinant 💦

Key Features

Host Species

Rabbit

MW • 17kD (Calculated) 17kD (Observed) Reactivity • Human, Mouse, Rat, Applications
• WB,IF,IP,ELISA

Recommended Dilution Ratios

WB 1:1000-1:5000 IF 1:200-1:1000 ELISA 1:5000-1:20000 IP 1:50-1:200

Storage

Storage* -15°C to -25°C/1 year(Do not lower than -25°C)

Formulation PBS, 50% glycerol, 0.05% Proclin 300, 0.05% BSA

Basic Information

Clonality	Monoclonal
Clone Number	PT0242R

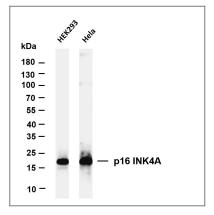
Immunogen Information

Specificity Endogenous

Target Information

Gene name	CDKN2A CDKN2 MTS1		
Protein Name	p16 INK		
	Organism	Gene ID	UniProt ID
	Human	<u>1029;</u>	<u>P42771;</u>
Cellular Localization	Cytoplasm, Nucleus		
Tissue specificity	Widely expressed but not detected in specific.	brain or skeletal muscle. Isof	orm 3 is pancreas-
Function	Widely expressed but not detected in brain or skeletal muscle. Isoform 3 is pancreas- specific. Alternative products:Isoform 1 and isoform 4 arise due to the use of two alternative first exons joined to a common exon 2 at the same acceptor site but in different reading frames, resulting in two completely different isoforms,Disease:Defects in CDKN2A are a cause of Li- Fraumeni syndrome (LFS) [MIM:151623]. LFS is a highly penetrant familial cancer phenotype usually associated with inherited mutations in TP53.,Disease:Defects in CDKN2A are involved in tumor formation in a wide range of tissues.,Disease:Defects in CDKN2A are because of cutaneous malignant melanoma 2 (CMM2) [MIM:155601]. Inheritance is autosomal dominant. Malignant melanoma 2 (CMM2) [MIM:155601]. Inheritance is autosomal dominant. Malignant melanoma 2 (CMM2) [MIM:156671], Inheritance is autosomal dominant. Malignant melanoma 3 (CMM2) [MIM:156761]. Inheritance is autosomal dominant. Malignant melanoma syndrome (FAMMMPC) [MIM:606719], Disease:Defects in CDKN2A are the cause of familial atypical multiple mole melanoma-pancreatic carcinoma syndrome (FAMMMPC) [MIM:606719], Disease:Defects in CDKN2A are the cause of melanoma-astrocytoma syndrome [MIM:15575]. The melanoma-astrocytoma syndrome is characterized by a dual predisposition to melanoma and neural system tumors, commonly astrocytoma., Function:As as a negative regulator of the proliferation of normal cells by interacting strongly with CDK4 and CDK6. This inhibits their ability to interact with cyclins D and to phosphorylate the retinoblastoma protein., Function:Capable of inducing cell cycle arrest in G1 and G2 phases. Acts as a tumor supressor. Binds to MDM2 and blocks its nucleocytoplasmic shuttling by sequestering it in the nucleolus. This inhibits the oncogenic action of MDM2 by blocking MDM2-induced degradation of p53 and enhancing p53- independent manner by preventing the activation of cyclin B1/CDC2 complexes. Binds to BCL6 and down-regulates BCL6-induced transcriptional repression. Binds t		

Validation Data



Various whole cell lysates were separated by 4-20% SDS-PAGE, and the membrane was blotted with anti-p16 INK4A antibody. The HRP-conjugated Goat anti-Rabbit IgG(H + L) antibody was used to detect the antibody. Lane 1: HEK293 Lane 2: Hela Predicted band size: 17kDa Observed band size: 17kDa

Contact information

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Please scan the QR code to access additional product information: p16 INK4A (PT0242R) PT® Rabbit mAb

For Research Use Only. Not for Use in Diagnostic Procedures.

Antibody | ELISA Kits | Protein | Reagents