

MIIP Rabbit pAb

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Catalog # AP58329

Product Information

Application	IHC-P, IHC-F, IF
Primary Accession	Q5JXC2
Reactivity	Human
Predicted	Mouse, Rat, Dog, Horse, Rabbit
Host	Rabbit
Clonality	Polyclonal
Calculated MW	42824
Physical State	Liquid
Immunogen	KLH conjugated synthetic peptide derived from human MIIP
Epitope Specificity	251-350/388
Isotype	IgG
Purity	affinity purified by Protein A

Buffer	0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.
SUBUNIT	Interacts with IGFBP2.
Post-translational modifications	Isoform 2 is degraded by the ubiquitin-proteasome pathway.
Important Note	

Background Descriptions	<p>This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.</p> <p>MIIP has 3 SEG (segments of low compositional complexity) domains, an RGD motif, and several potential phosphorylation sites. The C-terminal region of IGFBP2 interacts with a central 44-amino acid sequence of MIIP. MIIP inhibits glioma cells invasion and down-regulates adhesion- and motility-associated genes such as NFKB2 and ICAM1. It exhibits opposing effects to IGFBP2 on cell invasion. There are 2 named isoforms due to alternative splicing. Isoform 1 is expressed in brain but underexpressed in glioma tissues, at protein level. Isoform 2 is not detected in normal organs, but is expressed in gliomas with increasing levels with glioma progression. On the contrary, at protein level, isoform 2 is not detected in gliomas, suggesting that this isoform is unstable in glioma cells. Isoform 2 is degraded by the ubiquitin-proteasome pathway.</p>
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Additional Information

Gene ID	60672
Other Names	Migration and invasion-inhibitory protein, IGFBP2-binding protein, Invasion-inhibitory protein 45, Iip45, MIIP, IIP45
Target/Specificity	Ubiquitous. Isoform 1 is expressed in brain but underexpressed in glioma tissues, at protein level. Isoform 2 is not detected in normal organs, but is expressed in gliomas with increasing levels with glioma progression. On the contrary, at protein level, isoform 2 is not detected in gliomas, suggesting that

this isoform is unstable in glioma cells.

Dilution

IHC-P=1:100-500,IHC-F=1:100-500,IF=1:100-500

Storage

Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C.

Protein Information

Name

MIIP

Synonyms

IIP45

Function

Inhibits glioma cells invasion and down-regulates adhesion- and motility-associated genes such as NFKB2 and ICAM1. Exhibits opposing effects to IGFBP2 on cell invasion.

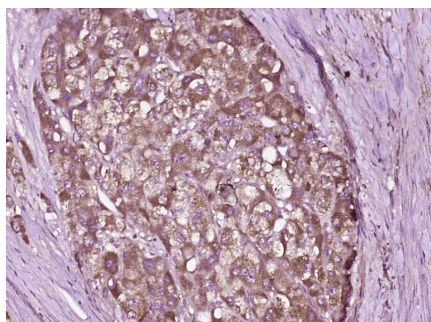
Tissue Location

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Background

MIIP has 3 SEG (segments of low compositional complexity) domains, an RGD motif, and several potential phosphorylation sites. The C-terminal region of IGFBP2 interacts with a central 44-amino acid sequence of MIIP. MIIP inhibits glioma cells invasion and down-regulates adhesion- and motility-associated genes such as NFKB2 and ICAM1. It exhibits opposing effects to IGFBP2 on cell invasion. There are 2 named isoforms due to alternative splicing. Isoform 1 is expressed in brain but underexpressed in glioma tissues, at protein level. Isoform 2 is not detected in normal organs, but is expressed in gliomas with increasing levels with glioma progression. On the contrary, at protein level, isoform 2 is not detected in gliomas, suggesting that this isoform is unstable in glioma cells. Isoform 2 is degraded by the ubiquitin-proteasome pathway.

Images



Paraformaldehyde-fixed, paraffin embedded (Human liver carcinoma); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (MIIP) Polyclonal Antibody, Unconjugated (AP58329) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.