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anti-SERPINB2 antibody (Internal Region)

2 Images

Overview



Publication



Go to Product page

Quantity:	100 μg
Target:	SERPINB2
Binding Specificity:	Internal Region
Reactivity:	Human, Monkey

Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This SERPINB2 antibody is un-conjugated

Application:	Western Blotting (WB), ELISA	
Application.	vvesterii blottiiig (vvb), EElo/(

Product Details

Product Details	
Immunogen:	This affinity purified antibody was prepared from whole rabbit serum produced by repeated immunizations with a peptide corresponding to an internal area of human PAI-2 protein. Immunogen Type: Peptide
Isotype:	IgG
Specificity:	This affinity-purified antibody is directed against human PAI-2 protein. The product was affinity purified from monospecific antiserum by immunoaffinity purification. Reactivity occurs against human PAI-2 protein. A BLAST analysis was used to suggest cross reactivity with PAI-2 proteins from several primates based on 94% homology with the immunizing sequence. Reactivity against homologues from other sources is not known.
Cross-Reactivity:	Monkey
Characteristics:	Extracellular plasminogen activator inhibitor type-2 (also known as PAI-2, Placental

plasminogen activator inhibitor, Monocyte Arg-serpin, Urokinase inhibitor, SERPINB2 serpin peptidase inhibitor, clade B (ovalbumin), member 2) is a coagulation factor and a potent inhibitor of urokinase-type plasminogen activator (u-PA) and also acts as a multifunctional protein. It is present in most cells, especially monocytes and macrophages. PAI-2 exists in two forms, a 60-kDa, secreted, extracellular, glycosylated form and a 43-kDa intracellular form. It is a multifunctional protein that plays a role in cell differentiation, in prevention of programmed cell death, in the regulation of cell proliferation, in the inhibition of microbial proteinases and in the protection against stromal degradation. High levels of the PAI-2 protein are associated with a good prognosis in breast cancer, small cell lung, ovarian cancer, and inhibition of metastasis. PAI-2 also plays a role in inflammation on the surface of the eye. PAI-2 may cooperate with pRb2/p130 in modulating PAI-2 gene expression by chromatin remodeling.

Purification:

affinity purified

Sterility:

Target:

Sterile filtered

SERPINB2

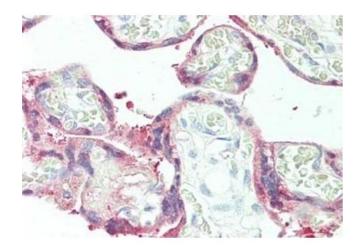
Target Details

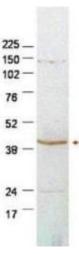
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Alternative Name:	Plasminogen Activator Inhibitor 2 (SERPINB2 Products)
Target Type:	Amino Acid
Background:	Extracellular plasminogen activator inhibitor type-2 (also known as PAI-2, Placental
	plasminogen activator inhibitor; Monocyte Arg-serpin; Urokinase inhibitor; SERPINB2 serpin
	peptidase inhibitor, clade B (ovalbumin), member 2) is a coagulation factor and a potent
	inhibitor of urokinase-type plasminogen activator (u-PA) and also acts as a multifunctional
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	forms, a 60-kDa, secreted, extracellular, glycosylated form and a 43-kDa intracellular form. It is a
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	death, in the regulation of cell proliferation, in the inhibition of microbial proteinases and in the
	protection against stromal degradation. High levels of the PAI-2 protein are associated with a
	good prognosis in breast cancer, small cell lung, ovarian cancer, and inhibition of metastasis.
	PAI-2 also plays a role in inflammation on the surface of the eye. PAI-2 may cooperate with
	pRb2/p130 in modulating PAI-2 gene expression by chromatin remodeling.
	Synonyms: PAI-2, Placental plasminogen activator inhibitor; Monocyte Arg-serpin; Urokinase
	inhibitor; SERPINB2 serpin peptidase inhibitor, clade B (ovalbumin), member 2
Gene ID:	5055, 4505595

Target Details UniProt: P05120 Pathways: Autophagy **Application Details** This affinity purified antibody has been tested for use in ELISA and western blot. Specific **Application Notes:** conditions for reactivity should be optimized by the end user. Expect a band 42-46.6 kDa in size corresponding to PAI-2 protein by western blotting in the appropriate cell lysate or extract. Comment: Gene Name: SERPINB2 Restrictions: For Research Use only Handling Format: Liquid Concentration: 1.2 mg/mL Buffer: 0.02 M Potassium Phosphate, 0.15 M Sodium Chloride, pH 7.2 Preservative: Sodium azide Precaution of Use: This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only. 4 °C/-20 °C Storage: Store vial at 4 °C prior to restoration. For extended storage aliquot contents and freeze at -20 °C Storage Comment: or below. Avoid cycles of freezing and thawing. Centrifuge product if not completely clear after standing at room temperature. This product is stable for several weeks at 4 °C as an undiluted liquid. Dilute only prior to immediate use. Expiration date is three (3) months from date of opening. 3 months **Expiry Date: Publications**

Product cited in:

Porchet, Probst, Dráberová, Dráber, Riederer, Riederer: "Differential subcellular localization of phosphorylated neurofilament and tau proteins in degenerating neurons of the human entorhinal cortex." in: **Neuroreport**, Vol. 14, Issue 7, pp. 929-33, (2003) (PubMed).





Immunohistochemistry (Paraffin-embedded Sections)

Image 1. Immunohistochemistry of rabbit anti-Plasminogen Activator 2 antibody (ABIN349627). Tissue: heart. Fixation: formalin fixed paraffin embedded. Antigen retrieval: not required. Primary antibody: Anti-Plasminogen Activator 2 at 10 μg/mL for 1 h at RT. Secondary antibody: Peroxidase rabbit secondary antibody at 1:10,000 for 45 min at RT. Staining: Plasminogen Activator II as precipitated red signal with hematoxylin purple nuclear counterstain.

Western Blotting

Image 2. Western blot using affinity purified anti-PAI-2 antibody shows detection of endogenous PAI-2 in 50 μg of HeLa whole cell lysates. The band at ~42 kDa (arrowhead) corresponds to PAI-2. Faint non-specific bands are also noted at lower and higher molecular weights positions. Primary antibody was used at a 1:500 dilution in 5% BLOTTO in PBS reacted overnight at 4°C. The membrane was washed and reacted with a 1:5,000 dilution of HRP-conjugated Gt-a-Rabbit IgG for 45 min at room temperature with ECL used for detection. Molecular weight estimation was made by comparison to prestained MW markers. Personal communication, Luca D'Agostino, Temple University, Philadelphia, PA