abcam

Product datasheet

Recombinant human PAI1 (mutated N150H + K154T + Q319L + M354I) protein ab92969

1 Image

Description

Product name Recombinant human PAI1 (mutated N150H + K154T + Q319L + M354I) protein

Biological activity >95% active by uPA titration. Kinetic Data: Second order rate constants for inhibition of: uPA =

 $5.1 \times 10^6 \,\mathrm{M}^{-1} \mathrm{s}^{-1} \,\mathrm{tPA} = 7.9 \times 10^5 \,\mathrm{M}^{-1} \mathrm{s}^{-1}$

Purity > 95 % SDS-PAGE.

> 99% pure by SDS PAGE

Expression system Escherichia coli

Protein length Full length protein

Animal free No

Nature Recombinant

Species Human

Specifications

Our Abpromise guarantee covers the use of ab92969 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications SDS-PAGE

Functional Studies

Form Liquid

Additional notes Solubility: > 2 mg/mL and < 5 mg/mL Ultraviolet: Absorbance (280nm) = 2.5 epsilon 0.1% = 1.0

ab92969 contains the following four mutations: K154T, Q319L, M354I and N150H. These mutations combine to confer great stability to the otherwise labile molecule essentially locking in

into the active conformation. It is an ideal choice for in vivo studies.

Preparation and Storage

Stability and Storage Shipped on Dry Ice. Store at -80°C.

pH: 6.60

Constituents: 0.82% Sodium phosphate, 0.0292% EDTA, 0.58% Sodium chloride

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General Info

Function This inhibitor acts as 'bait' for tissue plasminogen activator, urokinase, and protein C. Its rapid

interaction with TPA may function as a major control point in the regulation of fibrinolysis.

Tissue specificity Found in plasma and platelets and in endothelial, hepatoma and fibrosarcoma cells.

Involvement in diseaseDefects in SERPINE1 are the cause of plasminogen activator inhibitor-1 deficiency (PAI-1D)

[MIM:613329]. It is a hematologic disorder characterized by increased bleeding after trauma, injury, or surgery. Affected females have menorrhagia. The bleeding defect is due to increased fibrinolysis of fibrin blood clots due to deficiency of plasminogen activator inhibitor-1, which

inhibits tissue and urinary activators of plasminogen.

Note=High concentrations of SERPINE1 seem to contribute to the development of venous but not

arterial occlusions.

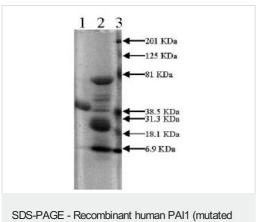
Sequence similarities Belongs to the serpin family.

Post-translational Inactivated by proteolytic attack of the urokinase-type (u-PA) and the tissue-type (TPA), cleaving modifications the 369-Arg-

the 369-Arg--Met-370 bond.

Cellular localization Secreted.

Images



SDS-PAGE - Recombinant human PAI1 (mutated N150H + K154T + Q319L + M354I) protein

(ab92969)

10% SDS Page Gelcode Blue Stain

- 1. 3ug ab92969 reduced
- 2. 3ug ab92969 + 12ug Urokinase reduced
- 3. prestained standard

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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