## SANTA CRUZ BIOTECHNOLOGY, INC.

# PAF acetylhydrolase (H-130): sc-33137



## BACKGROUND

The Platelet Activating Factor (PAF) Acetylhydrolases catalyze hydrolysis of the sn-2 ester bond of PAF and related pro-inflammatory phospholipids and thus attenuate their bioactivity. The family of PAF Acetylhydrolases include one secreted plasma isozyme and four intracellular proteins. The intra-cellular isozymes are distinguished by differences in their primary sequence, tissue localization, subunit composition, and substrate preferences. The most thoroughly characterized intracellular isoform, lb, contains two homologous (63% identity) 26 kDa catalytic subunits ( $\alpha$ 1 and  $\alpha$ 2), which harbor all the enzyme's activity, and a regulatory  $\beta$  subunit. The  $\alpha$  subunits readily associate with very high affinity to form homodimers, and this dimerization is essential for both stability and catalytic activity. The  $\beta$  subunit is a product of the LIS1 gene, mutations of which cause Miller-Dieker lissencephaly.

## REFERENCES

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- 3. Kuijpers, T.W., van den Berg, J.M., Tool, A.T. and Roos, D. 2001. The impact of platelet-activating factor (PAF)-like mediators on the functional activity of neutrophils: anti-inflammatory effects of human PAF-acetylhydrolase. Clin. Exp. Immunol. 123: 412-420.
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- 5. Quarck, R., De Geest, B., Stengel, D., Mertens, A., Lox, M., Theilmeier, G., Michiels, C., Raes, M., Bult, H., Collen, D., Van Veldhoven, P., Ninio, E. and Holvoet, P. 2001. Adenovirus-mediated gene transfer of human plateletactivating factor acetylhydrolase prevents injury-induced neointima formation and reduces spontaneous atherosclerosis in apolipoprotein E-deficient mice. Circulation 103: 2495-2500.

#### CHROMOSOMAL LOCATION

Genetic locus: PLA2G7 (human) mapping to 6p12; Pla2g7 (mouse) mapping to 17 C.

#### SOURCE

PAF acetylhydrolase (H-130) is a rabbit polyclonal antibody raised against amino acids 22-151 mapping near the N-terminus of PAF acetylhydrolase of human origin.

## PRODUCT

Each vial contains 200  $\mu$ g lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

## **APPLICATIONS**

PAF acetylhydrolase (H-130) is recommended for detection of PAF acetylhydrolase of human and, to a lesser extent, mouse and rat origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1–2 µg per 100–500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for PAF Acetylhydrolase siRNA (h): sc-39691.

## **RECOMMENDED SECONDARY REAGENTS**

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use goat anti-rabbit IgG-HRP: sc-2004 (dilution range: 1:2000-1:100,000) or Cruz Marker<sup>™</sup> compatible goat antirabbit IgG-HRP: sc-2030 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use goat anti-rabbit IgG-FITC: sc-2012 (dilution range: 1:100-1:400) or goat anti-rabbit IgG-TR: sc-2780 (dilution range: 1:100-1:400) with UltraCruz<sup>™</sup> Mounting Medium: sc-24941.

## **STORAGE**

Store at 4° C, \*\*DO NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

## **RESEARCH USE**

For research use only, not for use in diagnostic procedures.

#### PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.