biotechne[®] RDSYSTEMS

Human Matriptase/ST14 Catalytic Domain Antibody

Antigen Affinity-purified Polyclonal Sheep IgG Catalog Number: AF3946

DESCRIPTION			
Species Reactivity	Human		
Specificity	Detects human Matriptase/ST14 Catalytic Domain in direct ELISAs and Western blots.		
Source	Polyclonal Sheep IgG		
Purification	Antigen Affinity-purified		
Immunogen	<i>E. coli</i> -derived recombinant human Matriptase/ST14 Catalytic Domain Val615-Val855 Accession # Q9Y5Y6		
Formulation	Lyophilized from a 0.2 μm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied either lyophilized or as a 0.2 μm filtered solution in PBS		

APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.					
	Recommended Concentration	Sample			
Western Blot	0.1 μg/mL	Recombinant Human Matriptase/ST14 Catalytic Domain			
Intracellular Staining by Flow Cytometry	2.5 µg/10 ⁶ cells	See Below			
CyTOF-ready	Ready to be labeled using established conjugation methods. No BSA or other carrier proteins that could interfere with conjugation.				

DATA			
Intracellular Stain	ing by Flow Cytometry Detection of Matriptase/ST14 Catalytic Domain in PC-3 Human Cell Line by Flow Cytometry. PC-3 human prostate cancer cell line was stained with Sheep Artii-Human Matriptase/ST14 Catalytic Domain Antigen Affinity-purified Polycional Antibody (Catalog # AF3946, filled histogram), or control antibody (Catalog # 5-001-A, open histogram), orlowed by NorthernLights™ 557-conjugated Anti-Sheep IgG Secondary Antibody (Catalog # NL010). To facilitate intracellular staining, cells were fixed with paraformaldehyde and permeabilized with saponin.		
PREPARATION AND	STORAGE		
Reconstitution	Reconstitute at 0.2 mg/mL in sterile PBS.		
Shipping	hipping The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C		
Stability & Storage Use a manual defrost freezer and avoid repeated freeze-thaw cycles. 12 months from date of receipt, -20 to -70 °C as supplied. 1 month, 2 to 8 °C under sterile conditions after reconstitution. 6 months, -20 to -70 °C under sterile conditions after reconstitution.			

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BACKGROUND

RDSYSTEMS

Human matriptase, encoded by the ST14 (suppression of tumorogenicity 14) gene, is also known as tumor associated differentially expressed gene 15 protein/TADG-15), epithin, and membrane-type serine protease 1/MT-SP1 (1). Predicted to have a significant role in tumor biology, matriptase may be a novel target for anti-cancer therapy (2). However, expressed in most human epithelia, matriptase is also important in several physiological processes (1). For example, it activates prostasin to initiate a protease cascade that is essential for epidermal differentiation (3), and it converts a single-chain IGFBP-rp1 into the two-chain form (4).

Matriptase is a type II transmembrane serine protease with a complex modular structure (1). The 855 amino acid (aa) sequence of human matriptase consists of a cytoplasmic tail (aa 1-55), a transmembrane domain (aa 56-76), and an extracellular portion (aa 77-855). The latter contains the following domains: SEA (aa 86-201), two CUBs (aa 214-334 and 340-447), four LDLRAs (aa 452-486, 487-523, 524-560, and 566-603), and a serine protease (aa 615-855). The physiological activation of the single-chain zymogen requires the cleavage at the SEA domain within the ER or Golgi, association with HAI-1, which facilitates the transport of the protease to the cell surface, and auto-cleavage at QAR-V(615)VGG (1). The activated matriptase is inhibited by HAI-1, and the resulting HAI-1 complex can be shed from the cell surface (1). R&D Systems rhST14 corresponds to the catalytic domain, and is inhibited effectively by rhHAI-1 and rhHAI-2A (R&D Systems, Catalog # 1048-PI and 1106-PI).

References:

- 1. List, K. *et al.* (2006) Mol. Med. **12**:1.
- 2. Uhland, K. (2006) Cell. Mol. Life Sci. 63:2968
- 3. Netzel-Arnett, S. et al. (2006) J. Biol. Chem. 281:32941.
- 4. Ahmed, S. et al. (2006) FEBS J. 273:615.

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