SANTA CRUZ BIOTECHNOLOGY, INC.

MMP-26 (E-14): sc-27778



BACKGROUND

Metalloproteinases (MMPs) are a family of proteins that are involved in the breakdown of the extracellular matrix during normal cellular events, including reproduction, tissue remodeling and embryonic development. MMP-26 (matrix metallopeptidase-26), also known as endometase or matrilysin-2, is a 261 amino acid metalloproteinase that is secreted as an inactive protein and is activated upon cleavage by extracellular proteinases. Expressed specifically in the placenta and uterus, MMP-26 hydrolyzes (and subsequently degrades) a variety of proteins such as Fibrinogen, Fibronectin, Vitronectin and collagen type IV (COL4). MMP-26 binds zinc and calcium as cofactors and, unlike other MMP family members, lacks a conserved C-terminal domain. MMP-26 is widely expressed in a number of malignant tumor lines where it is thought to play an important role in tissue remodeling events that are associated with carcinogenesis.

REFERENCES

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- de Coignac, et al. 2000. Cloning of MMP-26. A novel matrilysin-like proteinase. Eur. J. Biochem. 267: 3323-3329.
- 4. Li, W., et al. 2004. Matrix metalloproteinase-26 is associated with estrogen-dependent malignancies and targets α 1-antitrypsin serpin. Cancer Res. 64: 8657-8665.
- Pilka, R. et al. 2004. Endometrial TIMP-4 mRNA is high at midcycle and in hyperplasia, but down-regulated in malignant tumours. Coordinated expression with MMP-26. Mol. Hum. Reprod. 10: 641-650.
- Bister, V., et al. 2005. Matrilysins-1 and -2 (MMP-7 and -26) and metalloelastase (MMP-12), unlike MMP-19, are up-regulated in necrotizing enterocolitis. J. Pediatr. Gastroenterol. Nutr. 40: 60-66.
- Lee, S., et al. 2006. Coordinated peak expression of MMP-26 and TIMP-4 in preinvasive human prostate tumor. Cell Res. 16: 750-758.
- Ahokas, K., et al. 2006. Matrix metalloproteinases 21 and 26 are differentially expressed in esophageal squamous cell cancer. Tumour Biol. 27: 133-141.
- Bister, V., et al. 2007. Increased expression of matrix metalloproteinases-21 and -26 and TIMP-4 in pancreatic adenocarcinoma. Mod. Pathol. 20: 1128-1140.

STORAGE

Store at 4° C, **D0 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.

SOURCE

MMP-26 (E-14) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the C-terminus of Matrix metalloproteinase-26 of human origin.

PRODUCT

Each vial contains 200 μg IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-27778 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

MMP-26 (E-14) is recommended for detection of precursor and mature MMP-26 of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), 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 MMP-26 siRNA (h): sc-106230, MMP-26 shRNA Plasmid (h): sc-106230-SH and MMP-26 shRNA (h) Lentiviral Particles: sc-106230-V.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (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) Immunofluores-cence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

PROTOCOLS

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