Human MMP14 Protein, hFc Tag Cat. No. PME100675



## **PRODUCT INFORMATION**

Target	MMP14
Synonyms	MMP-14;MMP-X1;MT-MMP;MT-MMP 1;MT1- MMP;MT1MMP;MTMMP1;WNCHRS
Description	Recombinant Human MMP14 with C-terminal human Fc tag
Delivery	In Stock
Uniprot ID	P50281
<b>Expression Host</b>	HEK293
Тад	C-Human Fc Tag
Molecular Characterization	MMP14(Tyr112-Ala541) hFc(Glu99-Ala330)
Molecular Weight	The protein has a predicted molecular mass of 75.3 kDa after removal of the signal peptide. The apparent molecular mass of MMP14-hFc is approximately 55-70 kDa due to glycosylation.
Purity	The purity of the protein is greater than 95% as determined by SDS-PAGE and Coomassie blue staining.
Formulation & Reconstitution	Lyophilized from sterile PBS, pH 7.4. Normally 5 % – 8% trehalose is added as protectants before lyophilization. Please see Certificate of Analysis for specific instructions of reconstitution.
Storage & Shipping	Store at -20°C to -80°C for 12 months in lyophilized form. After reconstitution, if not intended for use within a month, aliquot and store at -80°C (Avoid repeated freezing and thawing). Lyophilized proteins are shipped at ambient temperature.
Background	Proteins of the matrix metalloproteinase (MMP) family are involved in the breakdown of extracellular matrix in normal physiological processes, such as embryonic development, reproduction, and tissue remodeling, as well as in disease processes, such as arthritis and metastasis. Most MMP's are secreted as inactive proproteins which are activated when cleaved by extracellular proteinases. However, the protein encoded by this gene is a member of the membrane-type MMP (MT-MMP) subfamily; each member of this subfamily contains a potential transmembrane domain suggesting that these proteins are expressed at the cell surface rather than secreted. This protein activates MMP2 protein, and this activity may be involved in tumor invasion. [provided by RefSeq, Jul 2008]
Usage	Research use only

Email: info@dimabio.com Website: www.dimabio.com



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Figure 1. Human MMP14 Protein, hFc Tag on SDS-PAGE under reducing condition.

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