

DNAxPAb

Hard-to-Find
Antibody

MMP20 DNAxPab

Catalog # H00009313-W01P Size 200 ug

Specification

Product Description	Rabbit polyclonal antibody raised against a full-length human MMP20 DNA using DNAx™ Immune technology.
Technology	DNAx™ Immune
Immunogen	Full-length human DNA
Sequence	MKVLPAAGLAVFLIMALKFSTAAPSLVAASPRWNNYRLAQAYLDKYYTNKEGHQIGEMVARGSNMIRKIKELQAFFGLQVTGKLDQTTMNVIKKPRCGVPDVANYRLFPGEPKWKKNTLTYSKYTPSMSSVEVDKAVEMALQAWSSAVPLSFVRINSGEADIMISFENGHDGDSYPFDGPRGTLAHAFAPGEGLGGDTHFDNAEKWTMGNTGNLFTVAAHEFGHALGLAHSTDPSALMYPTYKYKNPYGFHLPKDDVKGIQALYGPRKVFLGKPTLPHAPHHKPSIPDLCDSSSSFDAVTMLGKELLLFKDRIFWRRQVHLRTGIRPSTITSSFPQLMSNVDAAYEVAERGTAFFKGPYHWRGFGMQGPPRTYDFGFPRHVQQIDAAYVLREPQKTLFFVGDEYYSYDERKRMKEDYPKNTEEEFSGVNGQIDAAYELNGYYFFSGPKTYKYDTEKEDVVSVKSSSWIGC
Host	Rabbit
Reactivity	Human
Purification	Protein A
Quality Control Testing	Antibody reactive against mammalian transfected lysate.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Applications

- Western Blot (Transfected lysate)

[Protocol Download](#)

- Immunofluorescence (Transfected cell)
- Flow Cytometry (Transfected cell)

Gene Info — MMP20

Entrez GeneID [9313](#)

GeneBank Accession# [NM_004771.3](#)

Protein Accession# [NP_004762.2](#)

Gene Name MMP20

Gene Alias MMP-20

Gene Description matrix metalloproteinase 20

Omim ID [204700 604629](#)

Gene Ontology [Hyperlink](#)

Gene Summary

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. The protein encoded by this gene degrades amelogenin, the major protein component of dental enamel matrix, and so the protein is thought to play a role in tooth enamel formation. A mutation in this gene, which alters the normal splice pattern and results in premature termination of the encoded protein, has been associated with amelogenesis imperfecta. This gene is part of a cluster of MMP genes that localizes to chromosome 11q22.3. [provided by RefSeq]

Other Designations enamel metalloproteinase|enamelysin|matrix metalloproteinase 20|matrix metalloproteinase 20 (enamelysin)

Disease

- [Cardiovascular Diseases](#)
- [Diabetes Mellitus](#)
- [Edema](#)
- [Tobacco Use Disorder](#)