


Product datasheet

Anti-ENAM antibody ab118134

2 Images

Overview

Product name	Anti-ENAM antibody
Description	Mouse monoclonal to ENAM
Tested applications	Suitable for: WB, ELISA, Sandwich ELISA
Species reactivity	Reacts with: Recombinant fragment Predicted to work with: Human 
Immunogen	Recombinant fragment, corresponding to amino acids 1043-1142 of Human ENAM with proprietary tag (NP_114095).

Properties

Form	Liquid
Storage instructions	Shipped at 4°C. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw cycles.
Storage buffer	pH: 7.20 Constituent: 99% PBS
Purity	Protein A purified
Clonality	Monoclonal
Isotype	IgG2a
Light chain type	kappa

Applications

Our [Abpromise guarantee](#) covers the use of **ab118134** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB		Use a concentration of 1 - 5 µg/ml. Predicted molecular weight: 129 kDa. This antibody has only been tested in WB against the recombinant fragment used as immunogen. We have no data on the detection of endogenous protein.

Application	Abreviews	Notes
ELISA		Use at an assay dependent dilution.
Sandwich ELISA		Use a concentration of 10 µg/ml.

Target

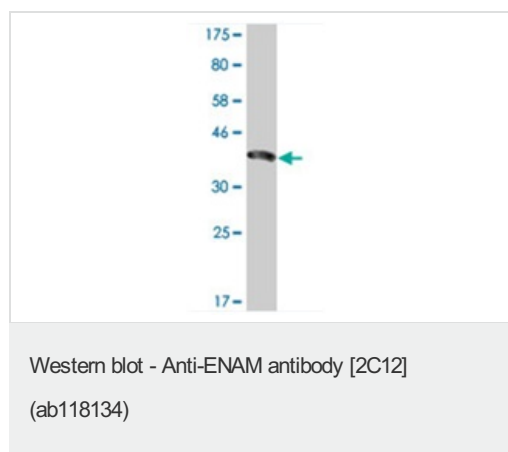
Function Involved in the mineralization and structural organization of enamel. Involved in the extension of enamel during the secretory stage of dental enamel formation.

Tissue specificity Expressed in tooth particularly in odontoblast, ameloblast and cementoblast.

Involvement in disease Defects in ENAM are the cause of amelogenesis imperfecta hypoplastic type 1B (AI1B) [MIM:104500]. AI1B is an autosomal dominant defect of enamel formation. Clinical manifestations may be variable. Some cases present with generalized enamel hypoplasia resulting in small, smooth, yellow and spaced teeth (smooth hypoplastic AI). Others show horizontal rows of pits, grooves or a hypoplastic area in the enamel (local hypoplastic AI). Defects in ENAM are the cause of amelogenesis imperfecta type 1C (AI1C) [MIM:204650]; also known as amelogenesis imperfecta hypoplastic with or without openbite malocclusion. AI1C is an autosomal recessive defect of dental enamel formation. Teeth show hypoplastic and unmineralized enamel, and a yellow-brown discoloration. Enamel defects can be associated with facial and oral features including vertical dysgnathia and anterior openbite malocclusion.

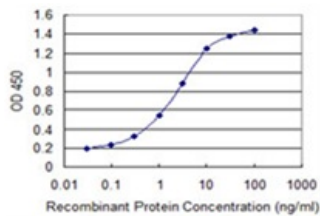
Cellular localization Secreted > extracellular space > extracellular matrix.

Images



Anti-ENAM antibody (ab118134) at 5 µg/ml + recombinant immunogen at 0.2 µg

Predicted band size : 129 kDa



Detection limit for recombinant tagged ENAM is 0.03 ng/ml as a capture antibody.

Sandwich ELISA - Anti-ENAM antibody [2C12]
(ab118134)

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