abcam

Product datasheet

Anti-Factor IX/PTC antibody ab128048

1 Image

Overview

Product name Anti-Factor IX/PTC antibody

Description Sheep polyclonal to Factor IX/PTC

Host species Sheep

Tested applications
Suitable for: IHC-P
Species reactivity
Reacts with: Human

Immunogen Full length native protein (purified) corresponding to Human Factor IX/PTC.

Positive control Human Liver tissue

General notes

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term.

Storage buffer pH: 7.40

Constituents: 0.24% HEPES, 0.88% Sodium chloride, 50% Glycerol (glycerin, glycerine)

Purity Immunogen affinity purified

Clonality Polyclonal

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab128048 in the following tested applications.

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

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Application	Abreviews	Notes
IHC-P		Use a concentration of 2.5 µg/ml.

Target

FunctionFactor IX is a vitamin K-dependent plasma protein that participates in the intrinsic pathway of blood coagulation by converting factor X to its active form in the presence of Ca(2+) ions, phospholipids, and factor VIIIa.

Tissue specificity Synthesized primarily in the liver and secreted in plasma.

Involvement in disease Defects in F9 are the cause of recessive X-linked hemophilia B (HEMB) [MIM:306900]; also

known as Christmas disease.

Note=Mutations in position 43 (Oxford-3, San Dimas) and 46 (Cambridge) prevents cleavage of the propeptide, mutation in position 93 (Alabama) probably fails to bind to cell membranes, mutation in position 191 (Chapel-Hill) or in position 226 (Nagoya OR Hilo) prevent cleavage of the

activation peptide.

Defects in F9 are the cause of thrombophilia due to factor IX defect (THR-FIX) [MIM:300807]. A

hemostatic disorder characterized by a tendency to thrombosis.

Sequence similaritiesBelongs to the peptidase S1 family.

Contains 2 EGF-like domains.

Contains 1 Gla (gamma-carboxy-glutamate) domain.

Contains 1 peptidase S1 domain.

DomainCalcium binds to the gamma-carboxyglutamic acid (Gla) residues and, with stronger affinity, to

another site, beyond the Gla domain.

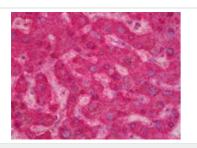
Post-translational Activated by factor Xla, which excises the activation peptide.

modifications The iron and 2-oxoglutarate dependent 3-hydroxylation of aspartate and asparagine is (R)

stereospecific within EGF domains.

Cellular localization Secreted.

Images



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Factor IX/PTC antibody (ab128048)

ab128048, at a concentration of 2.5 μ g/ml, staining Factor IX/PTC in paraffin-embedded Human liver tissue by Immunohistochemistry.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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