

Product datasheet for AM20168PU-N

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Factor H (CFH) Mouse Monoclonal Antibody [Clone ID: OX-24]

Product data:

Product Type: Primary Antibodies

Clone Name: OX-24

Applications: ELISA, IHC, IP, WB

Recommended Dilution: ELISA.

Immunohistochemistry on Paraffin Sections: 10 µg/ml.

Immunoprecipitation.

Western Blot.

Reactivity: Human
Host: Mouse
Isotype: IgG1

Clonality: Monoclonal

Immunogen: Purified Human complement factor H protein.

Specificity: Recognizes the Human serum complement protein factor H (155kD), binding to a different

epitope from OX23.

Recognizes a 43-49kD truncated form of factor H present at low level (1-5 µg/ml) in plasma

and urine.

In an Immunoprecipitation system, recognizes Factor H in other primates, but not in Bovine,

Sheep, Pig, Chicken or Rabbit sera.

This antibody inhibits the binding of factor H to surface bound C3b.

Recognizes Factor H in Western blotting procedures.

Formulation: State: Purified

State: Liquid purified Ig fraction containing 0.09% Sodium Azide as preservative.

Concentration: lot specific

Purification: Protein G Chromatography.

Conjugation: Unconjugated

Storage: Store the antibody undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer.

Avoid repeated freezing and thawing.

Stability: Shelf life: one year from despatch.





Gene Name: complement factor H

Database Link: Entrez Gene 3075 Human

P08603

Background: The complement Factor H protein is secreted into the bloodstream and acts in the regulation

of complement activation. Mutations leading to changes in this protein have been linked with HUS (hemolytic-uremic syndrome) and chronic hypocomplementemic nephropathy. Factor H is mainly synthesised in the liver but also in macrophages and endothelium. It is primarily aplasma glycoprotein but is also found in platelets and there is a membrane bound form on some leukocytes. Consisting of a single polypeptide, the major form of Factor H has a molecular weight of 155kDa. There are two truncated forms, a non-glycosylated 49 kDa form and a glycosylated 39-43 kDaform. Plasma concentrations are in the range 200-600mg/L for the 155 kDa form and 1-5mg/L for thetruncated forms. Factor H is a major regulatory protein of the complement system. By binding to C3b it either displacesor prevents the binding of Bb (activated Factor B). When bound to Factor H, C3b is susceptible tocleavage by Factor 1 to

yield iC3b.

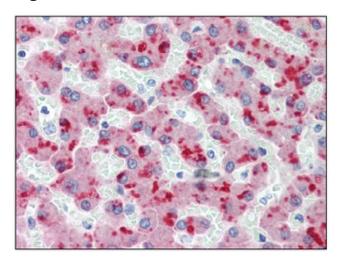
Factor H is released or modified following this cleavage. The regulatory role of Factor H is essential because C3bBb is not only a C5 convertase but a C3 convertaseand so has a positive

feedback effect, potentially consuming the entire C3 pool if unregulated.

Synonyms: CFH, HF, HF1, HF2, H factor 1

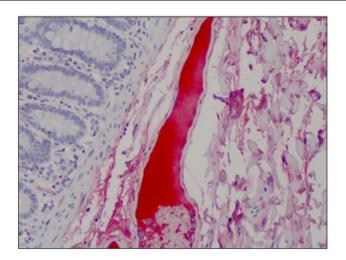
Protein Families: Druggable Genome, Secreted Protein
Protein Pathways: Complement and coagulation cascades

Product images:



Immunohistochemistry: AM20168PU-N Factor H antibody staining of Formalin-Fixed, Paraffin-Embedded Human Liver followed by biotinylated secondary antibody, alkaline phosphatase-streptavidin and chromogen.





Human Blood Plasma: Formalin-Fixed, Paraffin-Embedded (FFPE)