



## Monoclonal Antibodies: Anti-Human Factor H#3

For **Research Use Only**. Not for use in diagnostic procedures

### Background

Factor H is a fluid phase complement regulatory protein consisting of a single peptide chain of 20 short consensus repeat segments or CCP's with a molecular weight of approximately 155 KD.<sup>1</sup> Factor H regulates the alternative pathway of the complement system by modifying activity of the "feedback loop." It does this in three ways. First, it is a co-factor for the serine protease Factor I, which cleaves C3b to iC3b. iC3b has no hemolytic or amplification function, but may be bound by complement receptors. Second, Factor H prevents the formation of and accelerates the disassociation of the alternative pathway C3 convertase, C3bBb from cell surfaces. Finally, Factor H binds to polyanions on host cell surfaces and tissue matrices, such as basement membranes, blocking deposition of C3b. This later activity is leveraged by many pathogens as a mode of complement evasion.<sup>2</sup>

Recent studies have linked Factor H to hemolytic uremia syndrome (HUS),<sup>3</sup> age-related macular degeneration (AMD),<sup>4</sup> and membrano-proliferative glomerulonephritis. Factor H may also be elevated in certain cancers, including bladder cancer, potentially as a protective measure used by tumor cells to evade complement attack.

### Characterization

All of Quidel's monoclonal antibodies to complement antigens were prepared using intact complement proteins and are purified from mouse ascites fluid via protein A affinity chromatography. The prepared monoclonal antibodies are buffer exchanged in Borate Buffered Saline containing 0.02% NaN<sub>3</sub>.

The specificity of the Factor H (#3) (clone 131X) monoclonal antibody was established via a series of assays utilizing highly purified human Factor H and Factor H CCP fragments.<sup>4</sup> First, the antibody was shown by ELISA to bind to purified Factor H immobilized in microtiter wells. Free (unbound) Factor H and human serum but not other complement proteins were shown (via inhibition EIA) to inhibit the binding of this antibody to immobilized Factor H via Western Blot. This antibody was shown to bind specifically to CCP8-15a of Factor H. This antibody does not bind to Factor H Like Protein 1.

### Applications

Applications of this antibody have been evaluated by various research facilities, and include EIA,<sup>5</sup> Western Blot,<sup>4</sup> and FACS. Studies have shown that this antibody functionally inhibits Factor H and can be used to deplete Factor H from Human Serum.<sup>6,7</sup>

### Specifications

- Volume/vial: 100 µL
- Storage: 2°C to 8°C\* (≤ 30 days)
- Concentration: 1.0-1.2 mg/mL
- Buffer: Phosphate Buffered Saline
- Isotype: IgG2bk

### Species Cross Reactivity:

- Baboon, Cynomolgus Monkey

\*For long-term storage (> 30 days), aliquot and store at ≤ -20°C. Avoid repeated freeze-thaw.

### References

- <sup>1</sup>Pangburn, M.K. Differences between the binding sites of the complement regulatory proteins DAF, CR1 and Factor H on C3 Convertases. *J Immunol* 136:6 (1986).
- <sup>2</sup>Kraiczy, P., Würzner, R. Complement escape of human pathogenic bacteria by acquisition of complement regulators. *Mol Immunol* 43:21-44 (2006).
- <sup>3</sup>Atkinson, J.P., et al. Complement factor H and the hemolytic uremic syndrome. *JEM* Vol. 204, No. 6, June 11, 2007:1245-1248.
- <sup>4</sup>Sivaprasad, S. and Chong, N.V. The complement system and age related macular degeneration. *Eye* (2006), 1-6.

<sup>5</sup>Jokiranta, T.S., et al. Analysis of the recognition mechanism of the alternative pathway of complement by monoclonal anti-factor H antibodies: evidence for multiple interactions between H and surface bound C3b. *FEBS Ltr* 393(2-3):297-302 (1996).

<sup>6</sup>Cheng, Z., et al. Complement Factor H as a Marker for Detection of Bladder Cancer. *Clin Chem* 51:5 (2005).

<sup>7</sup>Junnikkala, S., et al. Exceptional resistance of Human H2 Glioblastoma Cells to Complement Mediated Killing by Expression and Utilization of Factor H and Factor H Like Protein I. *J Immunol* 164 (2000).

<sup>8</sup>On file at Quidel Corporation.

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