

#### OriGene Technologies, Inc.

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# Product datasheet for TA806031BM

## Factor I (CFI) Mouse Monoclonal Antibody (HRP conjugated) [Clone ID: OTI4A8]

## **Product data:**

| Product Type:           | Primary Antibodies   |
|-------------------------|--|
| Clone Name:             | OTI4A8   |
| Applications:           | IHC, WB  |
| Recommended Dilution:   | WB 1:2000  |
| Reactivity:             | Human  |
| Host:                   | Mouse  |
| lsotype:                | lgG1   |
| Clonality:              | Monoclonal   |
| Immunogen:              | Human recombinant protein fragment corresponding to amino acids 340-583 of human<br>CFI(NP_000195) produced in E.coli. |
| Formulation:            | PBS (pH 7.3) containing 1% BSA, 50% glycerol.  |
| Concentration:          | 0.5 mg/ml  |
| Purification:           | Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography<br>(protein A/G)           |
| Conjugation:            | HRP  |
| Storage:                | Store at -20°C as received.  |
| Stability:              | Stable for 12 months from date of receipt.   |
| Predicted Protein Size: | 63.4 kDa   |
| Gene Name:              | complement factor l  |
| Database Link:          | <u>NP_000195</u><br><u>Entrez Gene 3426 Human</u><br><u>P05156</u>   |



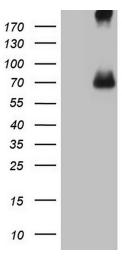
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### Sactor I (CFI) Mouse Monoclonal Antibody (HRP conjugated) [Clone ID: OTI4A8] – TA806031BM

Background: This gene encodes a serine proteinase that is essential for regulating the complement cascade. The encoded preproprotein is cleaved to produce both heavy and light chains, which are linked by disulfide bonds to form a heterodimeric glycoprotein. This heterodimer can cleave and inactivate the complement components C4b and C3b, and it prevents the assembly of the C3 and C5 convertase enzymes. Defects in this gene cause complement factor I deficiency, an autosomal recessive disease associated with a susceptibility to pyogenic infections. Mutations in this gene have been associated with a predisposition to atypical hemolytic uraemic syndrome, a disease characterized by acute renal failure, microangiopathic hemolytic anemia and thrombocytopenia. Primary glomerulonephritis with immmune deposits is another condition associated with mutation of this gene. [provided by RefSeq, Jul 2008]

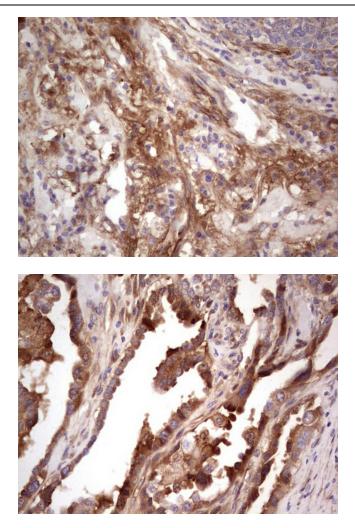
| Synonyms:         | AHUS3; ARMD13; C3b-INA; C3BINA; FI; IF; KAF  |
|-------------------|--|
| Protein Families: | Druggable Genome, Protease, Secreted Protein |
| Protein Pathways: | Complement and coagulation cascades          |

#### **Product images:**



HEK293T cells were transfected with the pCMV6-ENTRY control (Left lane) or pCMV6-ENTRY CFI ([RC216645], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-CFI. Positive lysates [LY424865] (100ug) and [LC424865] (20ug) can be purchased separately from OriGene.

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Immunohistochemical staining of paraffinembedded Carcinoma of Human lung tissue using anti-CFI mouse monoclonal antibody. ([TA806031]) Dilution: 1:150

Immunohistochemical staining of paraffinembedded Adenocarcinoma of Human ovary tissue using anti-CFI mouse monoclonal antibody. ([TA806031]) Dilution: 1:150

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