

## DESCRIPTION

<b>Species Reactivity</b>	Human
<b>Specificity</b>	Detects human RGM-C/Hemojuvelin in direct ELISAs.
<b>Source</b>	Monoclonal Mouse IgG <sub>2B</sub> Clone # 751741
<b>Purification</b>	Protein A or G purified from cell culture supernatant
<b>Immunogen</b>	Mouse myeloma cell line, NS0-derived human RGM-C/Hemojuvelin Gln36-Asp400 Accession # Q8ZVN8
<b>Formulation</b>	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied either lyophilized or as a 0.2 µm filtered solution in PBS.

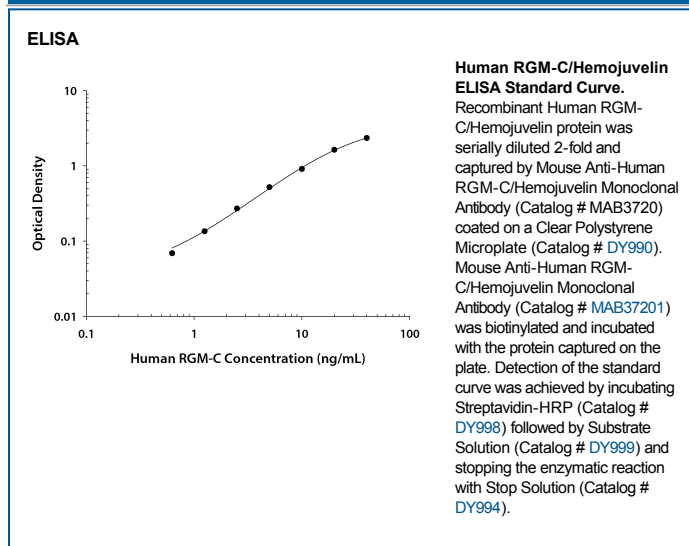
## APPLICATIONS

**Please Note:** Optimal dilutions should be determined by each laboratory for each application. *General Protocols* are available in the *Technical Information* section on our website.

**ELISA** This antibody functions as an ELISA capture antibody when paired with Mouse Anti-Human RGM-C/Hemojuvelin Monoclonal Antibody (Catalog # [MAB37201](#)).

*This product is intended for assay development on various assay platforms requiring antibody pairs. We recommend the Human RGM-C/Hemojuvelin DuoSet ELISA Kit (Catalog # [DY3720-05](#)) for convenient development of a sandwich ELISA.*

## DATA



## PREPARATION AND STORAGE

<b>Reconstitution</b>	Reconstitute at 0.5 mg/mL in sterile PBS.
<b>Shipping</b>	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C
<b>Stability &amp; Storage</b>	<b>Use a manual defrost freezer and avoid repeated freeze-thaw cycles.</b> <ul style="list-style-type: none"> <li>● 12 months from date of receipt, -20 to -70 °C as supplied.</li> <li>● 1 month, 2 to 8 °C under sterile conditions after reconstitution.</li> <li>● 6 months, -20 to -70 °C under sterile conditions after reconstitution.</li> </ul>

**BACKGROUND**

RGM-C, also known as hemojuvelin, is a member of the repulsive guidance molecule (RGM) family of GPI-linked neuronal and muscle membrane glycoproteins (1, 2). RGM-C is expressed in striated muscle and periportal hepatocytes (3 - 5). The protein undergoes partial cleavage intracellularly, resulting in a disulfide-linked dimer of the 14 kDa N-terminal and 33 kDa C-terminal portions (4, 6, 7). The N-terminal fragment contains an RGD motif, while the C-terminal fragment carries the GPI attachment site (4, 7). Two alternatively spliced isoforms lack either approximately half or the entire N-terminal fragment. Full length RGM-C can also be released from the cell and circulates in the blood (6, 8). RGM-C is disrupted in type 2A juvenile hemochromatosis, a hereditary iron homeostasis disorder characterized by excessive iron accumulation (5). In mouse, loss of RGM-C function results in decreased expression of the iron regulatory hormone hepcidin and increased iron deposition in liver, pancreas, and heart (5, 9). Membrane associated RGM-C upregulates hepcidin while soluble RGM-C downregulates hepcidin expression (8). This appears to be an iron-responsive regulatory system, as high blood iron levels reduce the amount of soluble RGM-C produced (8). RGM-C, similar to RGM-A, associates with neogenin (7). Disease-related point mutations can prevent internal RGM-C cleavage or its ability to interact with neogenin (6, 7). Experimental inflammatory conditions result in decreased RGM-C expression and increased hepcidin expression, although the two effects occur independently (5, 10). RGM-C also functions as a BMP coreceptor and enhances BMP-2 and BMP-4 signaling (11). In this context, RGM-C enhances the BMP-2 upregulation of hepatic hepcidin (11). Mature human RGM-C shares 89% amino acid (aa) sequence identity with mouse and rat RGM-C. It shares 49% and 44% aa sequence identity with human RGM-A and RGM-B, respectively.

**References:**

1. Papanikolaou, G. *et al.* (2004) *Nat. Genet.* **36**:77.
2. Schmidtmer, J. and D. Engelkamp (2004) *Gene Exp. Patterns* **4**:105.
3. Oldekamp, J. *et al.* (2004) *Gene Exp. Patterns* **4**:283.
4. Niederkofler, V. *et al.* (2004) *J. Neurosci.* **24**:808.
5. Niederkofler, V. *et al.* (2005) *J. Clin. Invest.* **115**:2180.
6. Kuninger, D. *et al.* (2006) *J. Cell Sci.* **119**:3273.
7. Zhang, A.S. *et al.* (2005) *J. Biol. Chem.* **280**:33885.
8. Lin, L. *et al.* (2005) *Blood* **106**:2884.
9. Huang, F.W. *et al.* (2005) *J. Clin. Invest.* **115**:2187.
10. Krijt, J. *et al.* (2004) *Blood* **104**:4308.
11. Babitt, J.L. *et al.* (2006) *Nat. Genet.* **38**:531.