

# Rat MAG/Siglec-4a Antibody

Monoclonal Mouse IgG<sub>2A</sub> Clone # 118423 Catalog Number: MAB538

DESCRIPTION			
Species Reactivity	Rat		
Specificity	Detects rat MAG/Siglec-4a in direct ELISAs and Western blots.		
Source	Monoclonal Mouse IgG <sub>2A</sub> Clone # 118423		
Purification	Protein A or G purified from hybridoma culture supernatant		
Immunogen	Mouse myeloma cell line NS0-derived recombinant rat MAG/Siglec-4a Gly20-Pro516 Accession # P07722		
Endotoxin Level	<0.10 EU per 1 µg of the antibody by the LAL method.		
Formulation	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.		

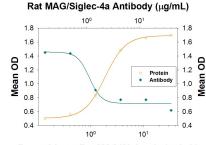
#### **APPLICATIONS**

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.

	Recommended Concentration	Sample
Western Blot	1 μg/mL	Recombinant Rat MAG/Siglec-4a Fc Chimera (Catalog # 538-MG)
Neutralization	Measured by its ability to neutralize MAG/Siglec-4a-mediated adhesion of human red blood cells. Kelm, S. et al. (1994) Current Biology 4:965. At 40 μg/mL, Mouse Anti-Rat MAG/Siglec-4a Monoclonal Antibody will neutralize approximately 60-85% of Recombinant Rat MAG/Siglec-4a Fc Chimera-dependent adhesion of human red blood cells.	

## DATA

## Neutralization



Recombinant Rat MAG/Siglec-4a (µg/mL)

# Cell Adhesion Mediated by MAG/Siglec-4a and Neutralization by Rat MAG/Sigle-4a Antibody. Recombinant Rat MAG/Siglec-4a Fc Chimera (Catalog # 538-MG), immobilized onto a microplate, supports the adhesion of human red blood cells in a dose-dependent manner (orange line). Adhesion elicited by

line). Adhesion elicited by Recombinant Rat MAG/Siglec-4a Fc Chimera (6 µg/mL) is neutralized (green line) by increasing concentrations of Mouse Anti-Rat MAG/Siglec-4a Monoclonal Antibody (Catalog # MAB538). At 40 µg/mL, this antibody will neutralize approximately 60-85% of Recombinant Rat MAG/Siglec-4a-dependent adhesion of human red blood cells.

### PREPARATION AND STORAGE

**Reconstitution** Reconstitute at 0.5 mg/mL in sterile PBS.

Shipping 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

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- 12 months from date of receipt, -20 to -70 °C as supplied.
- 1 month, 2 to 8 °C under sterile conditions after reconstitution.
- 6 months, -20 to -70 °C under sterile conditions after reconstitution.

Rev. 2/7/2018 Page 1 of 2





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#### BACKGROUND

MAG (Myelin-Associated Glycoprotein), a type I transmembrane glycoprotein containing five Ig-like domains in its extracellular domain is an adhesion molecule belonging to the immunoglobin superfamily. Within this superfamily, MAG, CD22, CD33, Schwann cell myelin protein, and sialoadhesin which bind specifically to cell-surface glycan containing sialic acid residues define the I-type sialyl lectin subgroup, also called the sialoadhesin family. Sialoadhesins mediate diverse biological processes through recognition of specific sialyted glycans on cell surface. MAG is expressed on myelinating oligodenrocytes and Schwann cells, and preferentially recognizes a2, 3-linked sialic acid on O-linked glycans and gangliosides. MAG exists as two isoforms which differ in the sequence and length of the cytoplasmic tail. The large form (71 kDa) and small form (67 kDa) arise from alternative spliced mRNAs. Although MAG might encounter haematopoietic cells and lymphocytes under pathologic conditions, it would normally be expected to interact with neuronal cells. It has been shown that MAG promotes axonal growth from neonatal DRG neurons and embryonic spinal neurons, but is a potent inhibitor of axonal re-growth from adult DRG and postnatal cerebellar neurons. MAG plays an important role in the interaction between axons and myelin. A soluble form of MAG containing the extracellular domain is released from myelin in large quantities and identified in normal human tissues and in tissues from patients with neurological disorders. This soluble MAG might contribute to the lack of CNS neuron regeneration after injury.

#### References:

- Kelm, S. et al. (1994) Current Biology 4:965.
- 2. McKerracher, L. et al. (1994) Neuron 13:805.
- 3. Tang, S. et al. (1997) Molecular and Cellular Neuroscience 9:333.
- 4. Cai, D. et al. (1999) Neuron 22:89.

