

Product Name:	Recombinant Mouse MOG Protein
Catalog Number:	AS-55150-100, AS-55150-500, AS-55150-1000
Lot Number:	See label on the vial
Amount/size:	100 μg, 500 μg, 1000 μg
Source:	The sequence (Accession # NP_034944) corresponding to the extracellular domain of mouse MOG along with a 6x His tag was expressed in <i>E. coli</i> . The recombinant mouse MOG (M-rMOG) was purified from urea denatured bacterial lysate using immobilized metal affinity chromatography (IMAC). The molecular weight of the recombinant mouse MOG is 14.2 kDa.
Activity:	Female C57BL/6NHsd and SJL/JCrHsd mice (9-10 weeks old) were immunized (s.c.) with 100 µg/animal of mouse rMOG in complete Freund's adjuvant followed by 400 ng/mouse injection of pertussis toxin on day 0 and day 2 (i.p.). Mice showed EAE symptoms such as limp tail, hind limb weakness, or hind limb paralysis after induction. Please note that no other EAE induction protocols were tested including IFA/cytokine model.
Purity:	Greater than 95% as determined by SDS-PAGE.
Endotoxin (EU/µg):	Less than 0.1 EU per 1 μ g of the protein as determined by Limulus Amebocyte Lysate (LAL) quantitative kinetic assay.
Storage:	The purified mouse rMOG is supplied as sterile and frozen at 1 mg/ml in 25 mM sodium acetate buffer (pH=4.0). Store at -80 °C, avoid repeated freeze-thaw cycles.

Instructions:

Myelin Oligodendrocyte Glycoprotein (MOG) is a member of the immunoglobulin superfamily and is expressed exclusively in central nervous system (CNS). Although MOG protein constitutes only 0.01-0.05% of the CNS myelin proteins, it was demonstrated that MOG protein is a crucial autoantigen for multiple sclerosis in humans and experimental autoimmune encephalomyelitis (EAE) in rodents and monkeys (1-5).

The purified mouse rMOG is recommended for in vitro studies such as T cell and B cell responses, cytokine response, antigen presentation, Western blotting, and ELISA as well as for in vivo study such as EAE induction in mice. The following dosages are recommended: 5-20 µg/ml for in vitro study and 50 µg per animal for in vivo study (1-5). *Please note, mouse MOG must be thoroughly mixed <u>directly</u> with Complete Freund's Adjuvant (CFA). Do not dilute recombinant mouse MOG with buffers that have pH greater than 4.5! Protein will precipitate at pH higher than 4.5!*



C57BL/6 Mice injected with Mouse MOG (1-125) Purified M-rMOG was gel at 3 µg/well and markers and purified CL=Crude Cell Lysa P=Purified M-rMOG. Figure 2. An Examp Five female C57/BL6 100 µg/animal mouse injection volume is 10 Subcutaneously (s.c. Pertussis Toxin (PT)

Figure 1. Mouse rMOG on SDS-PAGE. Purified M-rMOG was loaded onto 10-20% Tris-HCl gel at 3 μg/well and resolved at 200V for 60 minutes. Protein markers and purified M-rMOG (14.23 kDa) are indicated. CL=Crude Cell Lysate, FT=Flow Through, and

Figure 2. An Example of EAE Data Using Mouse rMOG. Five female C57/BL6 mice (9 weeks old) were injected with 100 µg/animal mouse rMOG (Cat. AS-55150) in CFA (total injection volume is 100 µl/site) at two flank sites subcutaneously (s.c.) along with 400 ng/animal of Pertussis Toxin (PT) on day 0 and 2. EAE scores may vary due to the animal health and housing conditions. This graph is for the reference only.

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Related Products

Product Name	Cat. #
Recombinant human MOG (1-125)	AS-55158
Recombinant rat MOG (1-125)	AS-55152
SensoLyte® Anti-Human MOG (1-125) Mouse IgG Specific ELISA Kit	AS-55153-M
SensoLyte® Anti-Human MOG (1-125) Rat IgG Specific ELISA Kit	AS-55153-R
SensoLyte® Anti-Human MOG (1-125) Human IgG Specific ELISA Kit	AS-55153-H
SensoLyte® Anti-Mouse MOG (1-125) IgG Quantitative ELISA Kit	AS-55156
SensoLyte® Anti-Rat MOG (1-125) IgG Quantitative ELISA Kit	AS-55157

References:

- 1. Jayaram Bettadapura et.al. (1998) Journal of Neurochemistry 70 (4): 1593-1599
- 2. Alfred R Oliver et al (2003) Journal of Immunology 171:462-468
- 3. Hans-Christian Von Budingen et.al. (2001) Journal of Clinical Immunology 21 (3): 155-170
- 4. Jerri-Anne Lyons et.al. (1999) European Journal of Immunology 29: 3432-3439
- 5. Hans-Christian Von Budingen et.al. (2004) European Journal of Immunology 34: 2072-2083

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