

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850 UNITED STATES Phone: +1-888-267-4436 Fax: +1-301-340-8606 techsupport@origene.com

OriGene Technologies GmbH

Schillerstr. 5 32052 Herford GERMANY Phone: +49-5221-34606-0 Fax: +49-5221-34606-11 info-de@origene.com

AM26335PU-N Monoclonal Antibody to MBL-A - Purified

Quantity:	0.1 mg
Concentration:	0.1 mg/ml
Background:	Mannose Binding Lectin (MBL), also called mannosebinding protein (MBP), is a calcium dependent oligomeric protein that belongs to the collectin family of proteins. It contains a collagen-like domain and a carbohydrate recognition domain enabling MBL to recognize carbohydrates (such as mannose and N-acetylglucosamine) on pathogens. MBL is able to activate the complement pathway independent of the classical and alternative complement activation pathways, by using attached mannose binding lectin-associated serine proteases (MASP-2) in an antibody- and C1q-independent manner. MASP-2 permits cleavage of C4 and C2 to form a C3 convertase. Once it has bound, MBL exhibits complement-dependent antibacterial activities such as microbial opsonization and/or microbial lysis via membrane attack complexes and therefore plays an important role in innate immunity. In human, MBL is encoded by a single gene, whereas in mice there are two homologous proteins, termed MBL-A and MBL-C. The MBL-A concentration in serum is about 6-fold lower compared to that of MBL-C MBL-A, but not MBL-C, was found to be an acute phase protein in casein and LPS-injection models. Moreover, it has been shown that MBL-A deficient mice have aberrant antigen-specific IgM responses and suffer from increased susceptibility to infection.
Uniprot ID:	<u>P39039</u>
NCBI:	<u>NP_034905.1</u>
GenelD:	<u>17194</u>
Host / Isotype:	Rat / IgG
Clone:	8G6
Immunogen:	Purified mouse MBL-A
Format:	State: Liquid 0.2 μm filtered Ig fraction Purification: Protein G Buffer System: PBS Preservatives: 0.02% sodium azide Stabilizers: 0.1% bovine serum albumin
Applications:	Immunohistochemistry on frozen sections (2,6): Tissue sections were fixed in 4% PBS- buffered formaldehyde and pretreated with 2% hydrogen peroxide in methanol for 20 minutes at 4°C to quench endogenous peroxidases. Primary antibody 8g6, 2µg/ml. As negative control rat IgG2a was used (Ref.2). The typical starting working dilution is 1:50. Immunoassays (2). Immunoflourescence (3,4). Western blot (1): A non-reduced sample treatment was used. The band sizes are 191, 263 and 316 kDa (Ref.1). The typical starting working dilution is 1:50.

For research and in vitro use only. Not for diagnostic or therapeutic work. Material Safety Datasheets are available at www.acris-antibodies.com or on request.

	ODICENIE
\sim	UNIGENE

	Other applications not tested. Optimal dilutions are dependent on conditions and should be determined by the user.
Specificity:	This antibody detects MBL-A.
Species Reactivity:	Tested: Mouse
Add. Information:	Note that the monoclonal antibody 8G6 is a calcium-dependent antibody.
General Readings:	 Liu H, Jensen L, Hansen S, Petersen SV, Takahashi K, Ezekowitz AB, et al. Characterization and quantification of mouse mannan-binding lectins (MBL-A and MBL-C) and study of acute phase responses. Scand J Immunol. 2001 May;53(5):489-97. PubMed PMID: 11309157. Windbichler M, Echtenacher B, Takahashi K, Ezekowitz RA, Schwaeble WJ, Jenseniuis JC, et al. Investigations on the involvement of the lectin pathway of complement activation in anaphylaxis. Int Arch Allergy Immunol. 2006;141(1):11-23. Epub 2006 Jun 23. PubMed PMID: 16804320. Held K, Thiel S, Loos M, Petry F. Increased susceptibility of complement factor B/C2 double knockout mice and mannan-binding lectin knockout mice to systemic infection with Candida albicans. Mol Immunol. 2008 Sep;45(15):3934-41. doi: 10.1016/j.molimm.2008.06.021. Epub 2008 Jul 30. PubMed PMID: 18672286. Petry F, Jakobi V, Wagner S, Tessema TS, Thiel S, Loos M. Binding and activation of human and mouse complement by Cryptosporidium parvum (Apicomplexa) and susceptibility of C1q- and MBL-deficient mice to infection. Mol Immunol. 2008 Jul;45(12):3392-400. doi: 10.1016/j.molimm.2008.04.010. Epub 2008 May 23. PubMed PMID: 18501966. Abe Y, Kuroda Y, Kuboki N, Matsushita M, Yokoyama N, Kojima N. Contribution of complement component C3 and complement receptor type 3 to carbohydrate- dependent uptake of oligomannose-coated liposomes by peritoneal macrophages. J Biochem. 2008 Nov;144(5):563-70. doi: 10.1093/jb/mvn101. Epub 2008 Aug 11. PubMed PMID: 18694897. Matthijsen RA, de Winther MP, Kuipers D, van der Made I, Weber C, Herias MV, et al. Macrophage-specific expression of mannose-binding lectin controls atherosclerosis in low-density lipoprotein receptor-deficient mice. Circulation. 2009 Apr 28;119(16):2188-95. doi: 10.1161/CIRCULATIONAHA.108.830661. Epub 2009 Apr 20.
	PubMed PMID: 19380618.
Pictures:	MBL-A (8G6) deposition in developing murine atherosclerotic lesions. Staining of frozen tissue sections with antibody 8G6 (AM26335PU-N). Anti-mouse MBL-A at 2µg/ml (2h, RT). MBL-A was detected on the intima to media border as well as throughout the media (insert). Furthermore, extensive MBL-A deposition was seen at sites of necrosis (upper right corner).

For research and in vitro use only. Not for diagnostic or therapeutic work. Material Safety Datasheets are available at www.acris-antibodies.com or on request.