

Human Complement Component C1r Antibody

Monoclonal Mouse IgG_{2B} Clone # 269104

Catalog Number: MAB1807

DESCRIPTION	
Species Reactivity	Human
Specificity	Detects human C1r in direct ELISAs and Western blots. In direct ELISAs, no cross-reactivity with recombinant human (rh) C1s or rhC1q R1 is observed. In Western blots, ths antibody recognizes rhC1r and the A/heavy chain.
Source	Monoclonal Mouse IgG _{2B} Clone # 269104
Purification	Protein A or G purified from hybridoma culture supernatant
Immunogen	Mouse myeloma cell line NS0-derived recombinant human Complement Component C1r Ser18-Asp705 Accession # NP_001724
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 Human Complement Component C1r (Catalog # 1807-SE) under non-reducing conditions only

 Immunoprecipitation
 25 μg/mL
 Conditioned cell culture medium spiked with Recombinant Human Complement

 Component C1r (Catalog # 1807-SE), see our available Western blot detection antibodies

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
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. 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.

BACKGROUND

The classical complement pathway plays a major role in innate immunity against infection. This pathway is triggered by C1, a multimolecular complex composed of the recognition protein C1q and two serine proteases, C1r and C1s. Following the C1q recognition, C1r is autoactivated, and in turn activates C1s, which cleaves C4 and C2, the C1 substrates (1). Both C1r and C1s activation involve cleavage of a specific Arg-lle bond, converting single-chain proenzymes into active proteases of disulfide bond-linked chains (A and B) (2). The A chains contain multiple domains in the order of CUB1-EGF-CUB2-CCP1-CCP2-Activation Peptide. The B chains contain the serine protease catalytic domain.

References:

- 1. Arlaud, G.J. et al. (2002) Biochem. Soc. Trans. 30:1001.
- 2. Lacroix, M. et al. (2001) J. Biol. Chem. 276:36233.
- 3. Journet A. and M. Tosi (1986) Biochem. J. 240:783

