

PRODUCTS FOR RESEARCH INTO THE FUNCTION AND DYSFUNCTION OF MITOCHONDRIAL COMPLEX II

MS201 COMPLEX II IMMUNOCAPTURE KIT

Isolates Complex II from human, mouse, rat and bovine tissues and cell lines

RESEARCH USES

The Complex II immunocapture kit allows isolation of the succinate dehydrogenase complex (E.C. 1.3.5.1) from small amounts of tissue. This facilitates subsequent analysis of assembly state, activity and the extent of post translational modifications including oxidative damage that occur with aging. Uses for the Complex II immunocapture kit include research on aging (1,2) genetic mitochondrial disease (3), and various cancers due to mutations in this enzyme complex such as paragangliomas and phaeo-chromocytomas (4,5).

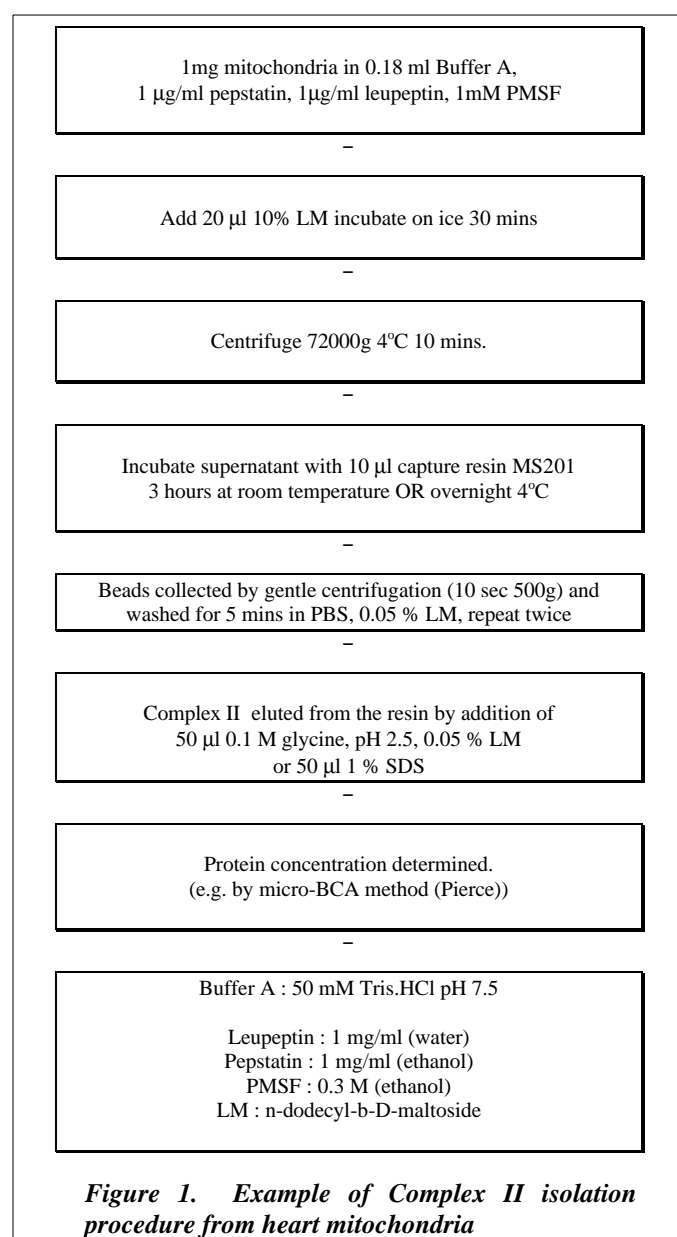
DESCRIPTION

The key component of the Complex II immunocapture kit is a monoclonal antibody able to selectively immunocapture the enzyme complex. The mAb is already covalently cross linked to Protein G-Agarose for convenience of use. This material is provided in batches of 25, 50 and 75 µl beads which have been charged with approximately 250, 500 and 750 µg of antibody respectively. When used as described in the protocol in Figure 1, 10 µl of beads are able to immunocapture approximately 5 µg of Complex II from heart mitochondria. Also provided are 2 mg of bovine heart mitochondria as a positive control to be used prior to, or during, isolation of Complex II from experimental samples. As an alternative, researchers can purchase the individual components i.e. 100 µg of mAb and 2 mg BHM (kit MS201c).

SUGGESTED PROTOCOL FOR IMMUNOCAPTURING COMPLEX II

The amount of Complex II that is captured in any experiment depends on both the amount of capture antibody and the amount of cell extract or isolated mitochondria used. Calculation of the amount of beads to be used in any experiment must also take account of the source of the material from which Complex II is to be isolated because mitochondria from different tissues have different concentrations of the enzyme complex. For example the levels of Complex II in mitochondria from cell culture material are around 10 fold less than in heart mitochondria. Figure 1 shows a schematic of a generic protocol developed for isolating Complex II from heart tissue.

Figure 2 shows the subunit structure of Complex II immunocaptured from beef heart and mouse heart mitochondria by Coomassie blue stained SDS-PAGE of immunoprecipitates obtained. The four subunits of the complex are clearly resolved. This antibody product is also reactive to and efficiently immunocaptures human Complex II.



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MATERIALS AND STORAGE

Kit MS201 contains the anti-Complex II immunocapture mAb covalently linked to protein G-agarose beads in 25, 50 or 75 μ l amounts. Beads have 8-10 μ g mAb bound per μ l bead volume. All volumes of bead resin are suspended in 400 μ l of PBS buffer (1.4 mM KH_2PO_4 , 8 mM Na_2HPO_4 , 140 mM NaCl, 2.7 mM KCl, pH 7.3) with 0.02 % sodium azide. Also included are 2 mg of purified bovine heart mitochondria resuspended in 400 μ l of heart mitochondria resuspension buffer (10 mM Tris.HCl, pH 7.8, 0.2 M sucrose, 0.2 mM EDTA, 1 mM PMSF). The antibody is shipped on ice. Upon receipt store the mAb at 4°C. The mitochondrial preparation should be aliquoted and stored at -20°C until use.

1. *J Biol Chem* (2003) 278, 22031-22036
2. *Mech Ageing Dev* (2004) 125, 41-46
3. *J Clin Invest* (2003) 111, 303-312
4. *J Pathol* (2003) 201, 480-486
5. *Clin Endocrinol (Oxf)* (2003) 59, 728-733

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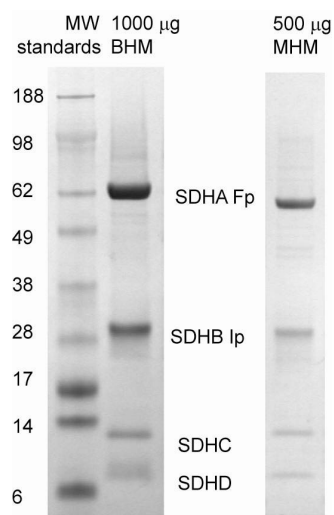


Figure 2. Immunocapture of Complex II bovine (BHM) and mouse heart mitochondria (MHM). Using 10 ml of capture resin MS201, the yield of Complex II from 1000 ng heart mitochondria is approximately 5 ng. However a yield detectable by Coomassie staining can be obtained from as little as 250 ng of total mitochondria. This yield is enough for most mass spectrometry applications.