abcam

Product datasheet

Complex I Immunocapture Kit ab109711

24 References 画像数 1

製品の概要

製品名

アッセイタイプ

種交差性

製品の概要

Complex I Immunocapture Kit

Quantitative

交差種: Mouse, Rat, Cow, Human

Optimized for isolation of Complex I from small amounts of tissue for activity assays and other analyses. Based on a highly specific antibody.

250 μg, 500 μg or 750 μg monoclonal antibody irreversibly crosslinked to protein G-agarose beads which can immunocapture \sim 60 μg, \sim 120 μg or \sim 180 μg respectively of Complex I from heart mitochondria.

ab109711 allows for isolation of Complex I from small amounts of tissue. This facilitates subsequent analysis of assembly state and activity. Thus the enzyme retains NADH-ferricyanide, NADH-CoQ1 and NADH hexaaminoruthenium reductase activities after isolation, and with added lipids it also shows significant rotenone sensitivity. Finally, the extent of post translational modifications including oxidative damage can be readily analyzed by proteomic approaches or antibody detection of these modifications. Uses for ab109711 include but are not limited to examining alterations of Complex I subunits in inherited mitochondrial diseases, Parkinson's disease, Alzheimer's disease, ALS, schizophrenia, and aging.

Note: The immunocapture protocol for this kit requires Abcam detergent (lauryl maltoside, **ab109857**).

Product supplied as a 6.25% slurry. For the 250 μ g, this consists of 250 μ g antibody conjugated to 25 μ L beads in a volume of 250 μ L.

Related products

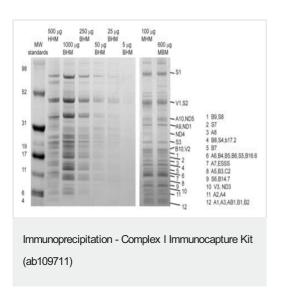
Review the <u>mitochondrial assay guide</u>, or the full <u>metabolism assay guide</u> to learn about more assays for metabolites, metabolic enzymes, mitochondrial function, and oxidative stress, and also how to assay metabolic function in live cells using your plate reader.

製品の特性

特記事項

内容	250 μg
Complex 1 monoclonal antibody coupled agarose beads	2 x 250µg

画像



Complex I was immunopurified from mitochondria isolated from human heart (HHM), cow/bovine heart (BHM), mouse heart (MHM) and mouse brain (MBM). The lanes were stained with Coomassie Brilliant Blue R. Bands were excised from the gel and proteolytically digested for mass spectrometry analysis."

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