abcam

Product datasheet

ATP synthase Enzyme Activity Microplate Assay Kit ab109714

★★★★★ 1 Abreviews 40 References 画像数 4

製品の概要

製品名 ATP synthase Enzyme Activity Microplate Assay Kit

検出方法 Colorimetric

サンプルの種類 Cell culture extracts, Tissue, Suspension cells, Tissue Extracts, Cell Lysate, Purified mitochondria

アッセイタイプ Enzyme activity

全工程の試験時間 6h 00m

種交差性 交差種: Rat, Cow, Human

非交差種: Mouse

製品の概要 ATP synthase Enzyme Activity Microplate Assay Kit ab109714 is used to determine the activity of

ATP synthase (Complex V) in a human or rat sample.

The ATP synthase enzyme is immunocaptured within the wells of the microplate and the enzyme activity is measured by monitoring the decrease in absorbance at 340 nm. The conversion of ATP to ADP by ATP synthase is coupled to the oxidation reaction of NADH to NAD+ with a reduction in

absorbance at 340 nm.

試験プラットフォーム Microplate reader

製品の特性

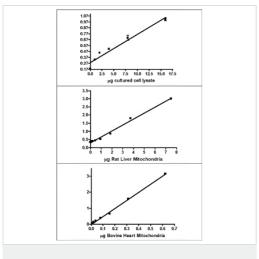
保存方法 Please refer to protocols.

内容	ラベル	96 tests
ATP synthase precoated Microtiter plate		1 unit
Buffer	Tube 1	1 x 10ml
Detergent		1 x 1ml

1

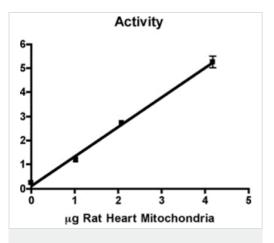
内容	ラベル	96 tests
Phospholipids		1 x 6ml
Reagent Mix		1 x 20ml

画像



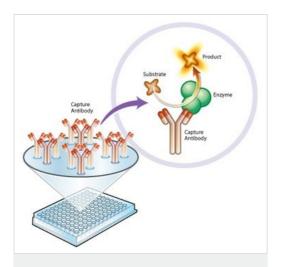
Examples of activity/load relationships for cultured cell lysate, rat liver mitochondria, and bovine heart mitochondria.

Functional Studies - ATP synthase Enzyme Activity Microplate Assay Kit (ab109714)



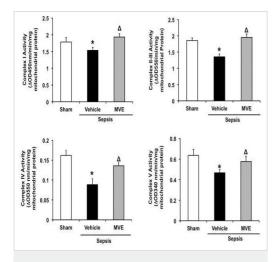
Functional study using ATP synthase Enzyme Activity Microplate Assay Kit (ab109714).

Functional Studies - ATP synthase Enzyme Activity Microplate Assay Kit (ab109714)



Principle of ATP synthase Enzyme Activity Microplate Assay Kit (ab109714).

Functional Studies - ATP synthase Enzyme Activity Microplate Assay Kit (ab109714)



Mitochondrial ROS dependent functional deficiency and structural impairment in cardiac mitochondria after sepsis.

Image courtesy of Yao X et al.PLoS One. 2015; 10(10): e0139416. doi: 10.1371/journal.pone.0139416.

Mitchondiral fractions from the heart tissue of Rats infected by S. pneumoniae, or given PBS sham control, were subjected to measurements of complex I-V activities. Complex I was measured with ab109721 (top left), Complex II + III were measured using ab109905 (top right), Complex IV was measured using ab109911 (bottom left) and Complex V was measured using ab109714 (bottom right).

Freshly isolated mitochondrial pellets were resuspened in PBS supplemented with 10% detergent provided in the kits. Protein concentrations of these mitochondrial lysates were estimated and 25 μ g (for complex I, IV and V) or 100 μ g (for complex II+III) mitochondrial protein was used per reaction. Enzyme activities were measured spectrophotometricly in triplicate and expressed as changes of absorbance per minute per mg protein.

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