abcam

Product datasheet

ATP Assay Kit (Colorimetric/Fluorometric) ab83355

8 Images

Overview

Notes

Product name ATP Assay Kit (Colorimetric/Fluorometric)

Detection method Colorimetric/Fluorometric

Sample type Urine, Serum, Plasma, Other biological fluids, Tissue Extracts, Cell Lysate

Assay type Quantitative

Sensitivity $< 1 \mu M$

Assay time 1h 00m

Product overview ATP Assay Kit (Colorimetric/Fluorometric) (ab83355) use a robust, simple method; the ATP

> assay protocol relies on the phosphorylation of glycerol to generate a product that is easily quantified by colorimetric (ODmax = 570 nm) or fluorometric (Ex/Em = 535/587 nm) methods.

This kit can detect as low as 1 μ M of ATP in various samples.

ATP assay protocol summary:

- add samples (deproteinized) and standards to wells
- add reaction mix and incubate for 30 min at room temp
- analyze with microplate reader

Chinese protocol available. See protocols section below.

If you require a more sensitive product, we recommend Luminescent ATP Detection Assay Kit

(ab113849), which can detect as low as 1 pM of ATP.

This product is manufactured by BioVision, an Abcam company and was previously called K354 ATP Colorimetric/Fluorometric Assay Kit. K354-100 is the same size as the 100 test size of

ab83355.

Related assays

Review the **cell health assay guide** to learn about kits to perform a **cell viability** assay, cytotoxicity assay and cell proliferation assay.

Review the metabolism assay guide to learn about assays for metabolites, metabolic enzymes, mitochondrial function, and oxidative stress, and also about how to assay metabolic function in

live cells using your plate reader.

How other researchers have used ATP Assay Kit ab83355

This ATP assay kit has been used in publications in a variety of sample types, including:

- Human: cell culture lysates¹, primary monocyte cell culture lysates², HCT116 cell culture supernatants³
- Mouse: heart tissue 4 , liver 5 , C2C12 and L929 cell lysates 6 , primary thymocyte cell culture lysates 7 , cardiac tissue 8
- Rat: primary hippocampal neuron cell culture lysates⁹, liver tissue¹⁰, skeletal muscle¹¹
- Pig: kidney cell culture lysates¹², heart tissue¹³
- C elegans tissue¹⁴
- Chlamydomonas reinhardtii algae¹⁵

References: 1 - Civallero Met al 2017, Na JY et al 2018, 2Gkirtzimanaki K et al 2018, 3Yang et al 2018; 4 - Singh SP et al 2018; 5 - Han SJ et al 2018; 6 - Alhindi Y et al 2019, Gregorczyk KP et al 2018; 7 - Simula L et al 2018; 8 - Litt MJ et al 2017, Samokhvalov V et al 2018; 9 - Zhao X et al 2018; 10 - Jing R et al 2018; 11 - Trinchese G et al 2018; 12 - Zou X et al 2018; 13 - Yuan F et al 2018; 14 - Pandey et al 2018; 15 - Ramanan R et al 2018

Platform

Microplate reader

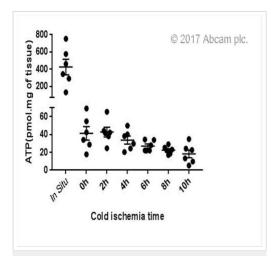
Properties

Storage instructions

Store at -20°C. Please refer to protocols.

Components	100 tests	2000 tests
Assay Buffer XXIII	1 x 25ml	20 x 25ml
ATP Standard	1 vial	20 vials
Converting Enzyme I	1 vial	20 vials
Developer IV	1 vial	20 vials
OxiRed™ Probe	1 x 200µl	20 x 200µl

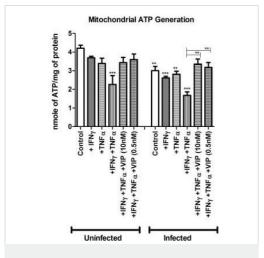
Images



ATP levels in Pancreatic Islet

Image courtesy of Mrs. Fotini Mouth

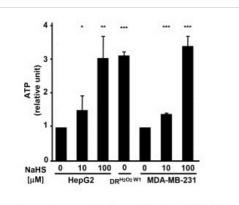
Ab83355 was used to determin ATP levels in rat pancreas islets as an ischemic marker to predict transplantation outcomes. We extracted ATP from fresh pancreas that have undergo different time of cold ischemia: 0, 2, 4, 6, 8 and 10h and in situ. ATP were extracted in Perchloric acid (PCA-2M) and grind using a Polytron. PCA were removed using potassium hydroxide (KOH – 2M) and pH was adjust around 7-8. Samples were conserved at -80°C before utilization.



ATP assay used to study mitochondrial disfunction after C. rodentium infection in mice

Image courtesy of Maiti AK et al. PLoS One. 2018; 13(9): e0204567. doi: 10.1371/journal.pone.0204567

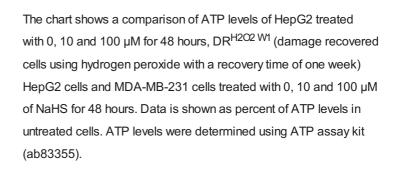
Maiti AK et al (2018) used ATP assay kit ab83355 to measure mitochondrial ATP generation in an in vitro mouse intestinal model treated with cytokines in the presence and absence of VIP (vasoactive intestinal peptide). VIP was induced by *C. rodentium* infection and cytokines.

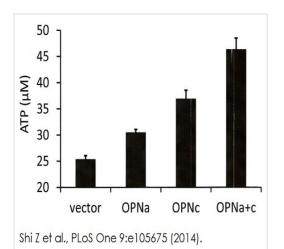


Sanokawa-Akakura R et al., PLoS One 9:e108537 (2014).

ATP assay performed with ab83355

Sanokawa-Akakura R et al., PLoS One 9(9), fig4b. doi: 10.1371/journal.pone.0108537 Reproduced under the Creative Commons license http://creativecommons.org/licenses/by/4.0/

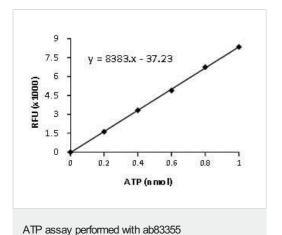




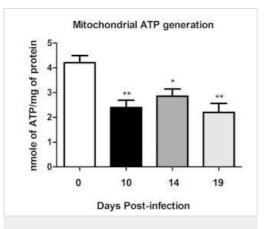
ATP assay performed with ab83355

Image from Shi Z et al., PLoS One 9(8), Fig 2 doi: 10.1371/journal.pone.0105675. Reproduced under the Creative Commons license http://creativecommons.org/licenses/by/4.0/

MCF-7 cells are transfected with vector, osteopontin-a, osteopontin-c or osteopontin -a plus -c. Cells are plated on poly-HEMA and seeded at 4 x 10^5 cells per well and incubated for two days under standard culture conditions. ATP levels are measured using ATP assay kit (ab83355).



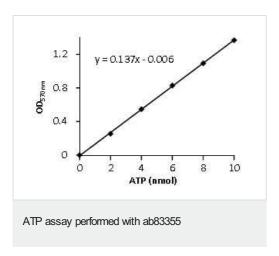
Example of fluorometric ATP assay standard curve.



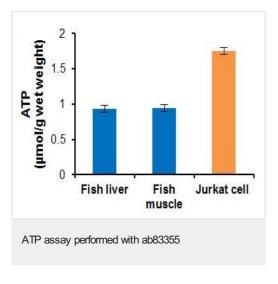
Maiti AK et al. (2015) used ATP assay kit <u>ab113852</u> to measure mitochondrial ATP generation in murine distal colon after *C.* rodentium infection.

ATP levels measured in mouse colon tissue

Image courtesy of Maiti A K et al. Sci Rep. 2015; 5: 1543. doi: 10.1038/srep15434. Reproduced under the Creative Commons License http://creativecommons.org/licenses/by/4.0/



Example of colorimetric ATP assay standard curve.



Quantitation of ATP in fish liver (2.5 μ l of 10 times diluted sample), fish muscle (5 μ l of 10 times diluted sample) and Jurkat cell lysate (5 ul) using fluorometric assay. Samples were spiked with known amounts of ATP (300pmol).

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