abcam

Product datasheet

ADP/ATP Ratio Assay Kit (Bioluminescent) ab65313

★★★★★ 2 Abreviews 108 References 画像数3

製品の概要

製品名 ADP/ATP Ratio Assay Kit (Bioluminescent)

検出方法 Luminescent

サンプルの種類 Tissue, Adherent cells, Suspension cells

アッセイタイプ Semi-quantitative

検出感度 < 100 cells/well

全工程の試験時間 0h 30m

ADP/ATP Ratio Assay Kit (Bioluminescent) ab65313 is based on the bioluminescent detection of ADP and ATP levels. It can be used for a rapid screening of apoptosis, necrosis, growth arrest,

and cell proliferation simultaneously in mammalian cells.

In the ADP/ATP assay protocol, luciferase catalyzes the conversion of ATP and luciferin to light, which in turn can be measured using a luminometer or Beta Counter. The ADP level is measured by its conversion to ATP that is subsequently detected using the same reaction.

The assay can be fully automatic for high throughput and is highly sensitive (detects 100 mammalian cells/well).

ADP/ATP Ratio assay protocol summary:

- transfer suspension cells, or nucleotide releasing buffer treated adherent cells, to plate
- add ATP reaction mix and incubate for 2 min
- analyze with luminescence plate reader to measure ATP
- after preparing ADP reaction mix and measuring luminescence levels again, add ADP reaction mix to same wells and incubate for 2 min
- analyze with luminescence plate reader to measure ADP

This product is manufactured by BioVision, an Abcam company and was previously called K255 ADP/ATP Ratio Bioluminescence Assay Kit, ApoSENSOR. K255-200 is the same size as the 200 test size of ab65313.

Changes in the ADP/ATP ratio have been used to differentiate the different modes of cell death and viability. Increased levels of ATP and decreased levels of ADP have been recognized in proliferating cells. In contrast, decreased levels of ATP and increased levels of ADP are recognized in apoptotic cells. The decrease in ATP and increase in ADP are much more pronounced in necrosis than apoptosis.

製品の概要

特記事項

Related assays

Review the <u>cell health assay guide</u> to learn about kits to perform a <u>cell viability</u> <u>assay</u>, <u>cytotoxicity assay</u> and <u>cell proliferation assay</u>.

Review the <u>metabolism assay guide</u> 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.

試験プラットフォーム

Microplate reader

製品の特性

保存方法

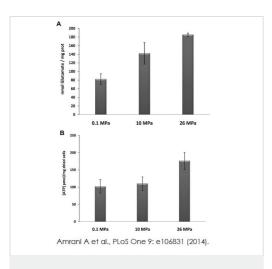
Store at -20°C. Please refer to protocols.

内容	200 tests
1X Reaction Buffer I	1 x 50ml
ADP Converting Enzyme	1 vial
ATP Monitoring Enzyme	1 vial
Enzyme Reconstitution Buffer	1 x 2ml

関連性

The changes in ADP/ATP ratio have been used to differentiate the different modes of cell death and viability. Increased levels of ATP and decreased levels of ADP have been recognized in proliferating cells. In contrast, decreased levels of ATP and increased levels of ADP are recognized in apoptotic cells. The decrease in ATP and increase in ADP are much more pronounced in necrosis than apoptosis.

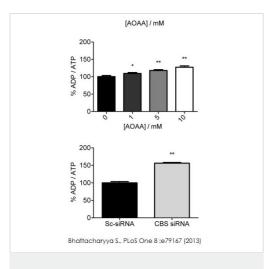
画像



Functional studies - ab65313

Image from Amrani Aet al., PLoS One 9(9). Fig 5A doi: 10.1371/journal.pone.0106831 Reproduced under the Creative Commons license http://creativecommons.org/licenses/by/4.0/

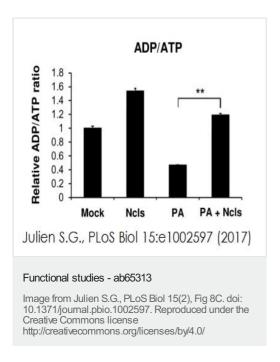
Quantitation of glutamate levels (A) using <u>ab138883</u> and intracellular ATP (B) using ab65313 in *D. hydrothermalis* cells grown under different pressure conditions.



ADP/ATP ratio in cystathionine-beta-synthase slienced A2780 (Ovarian cancer cell line) cells and AOAA (aminooxyacetic acid) treated A2780 cells were measured using ADP/ATP ratio assay kit (ab65313).

Functional studies - ab65313

Bhattacharyya S.,PLoS One 8(11), Fig 5d & f. doi: 10.1371/journal.pone.0079167 Reproduced under the Creative Commons license http://creativecommons.org/licenses/by/4.0/



ADP/ATP ratio were examined in both narciclasine (ncls) and vehicle (veh) treated C2C12 myotubes with or without PA treatment using ADP/ATP ratio assay kit (ab65313).

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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