abcam

Product datasheet

Cell Viability Assay Kit (Fluorometric - Green) ab112122

1 References 1 图像

概述

产品名称 Cell Viability Assay试剂盒(Fluorometric - Green)

检测方法 Fluorescent

样品类型 Adherent cells, Suspension cells

检测类型 Quantitative

种属反应性 与反应: Mammals, Other species

产**品概述** There are

There are a variety of parameters that can be used to monitor cell viability. The green fluorescent dye used in the kit is a hydrophobic compound. It easily permeates intact live cells and gets enhanced fluorescence upon entering into live cells. The hydrolysis of the non-fluorescent substrate by intracellular esterases generates a strongly green fluorescent hydrophilic product that is well-retained in the cell cytoplasm. The green fluorophore generated by the non-fluorescent substrate used in the kit has the spectral properties of fluorescein at Ex/Em = ~490 nm/520 nm. When well excited with the Argon Laser at 488 nm, the fluorophore emits intense green fluorescence at ~520 nm.

ab112122 provides all the essential components with an optimized cell-labeling protocol for fluorescence microplate assays. It can also be used with a fluorescence microscope equipped with a FITC filter set.

ab112122 provides an effective tool of labeling cells for fluorescence microplate and microscopic investigations of cellular functions. It is useful for a variety of studies, including cell adhesion, chemotaxis, multidrug resistance, cell viability, apoptosis and cytotoxicity. ab112122 is suitable for proliferating and non-proliferating cells.

Visit our **FAQs page** for tips and troubleshooting.

Review the **cell health assay guide** to learn about more kits to perform a **cell viability**

assay, cytotoxicity assay and cell proliferation assay.

平台 Microplate reader, Fluor. microscope, Flow cyt.

性能

说明

存放说明 Store at -20°C. Please refer to protocols.

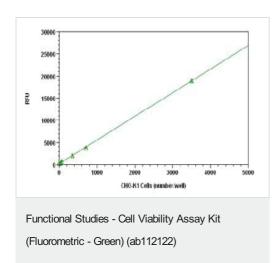
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组 件	5 x 96 tests
Assay Buffer	1 x 50ml
CellGreen fluorescent dye	5 vials
DMSO	1 x 200µl

相关性

Cell viability is a determination of living or dead cells, based on a total cell population. Cell viability assess healthy cells in a sample, with no distinction between dividing or quiescent cells. An increase in cell viability indicates cell growth, while a decrease in viability can generally be interpreted as the result of either toxic effects of compounds/agents or suboptimal culture conditions.

图片



CHO-K1 cell number response was measured with ab112122. CHO-K1 cells at 0 to 5,000 cells/well/100 μ L were seeded overnight in a black wall/clear bottom 96-well plate. The cells were incubated with 100 μ L/well of Green dye-loading solution for 1 hour at 37 °C. The fluorescence intensity was measured at Ex/Em = 490/ 525 nm. The fluorescence intensity was linear (R² = 1) to the cell number as indicated. The detection limit was 30 cells/well (n=6).

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