abcam

Product datasheet

10X RIPA Buffer ab156034

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Overview

Product name 10X RIPA Buffer

Tested applications Suitable for: WB, ELISA, SDS-PAGE, IP

General notes Abcam's 10X RIPA lysis buffer is an efficient means of cell lysis and protein solubilization for both

adherent and suspension cultured mammalian cells. This reagent effectively extracts cytoplasmic, nuclear and membrane proteins. It is compatible with many downstream applications, including

SDS-PAGE, Western blot, immunoprecipitation, ELISA and BCA assays.

Preparation: Dilute to 1X in deionized water

Abcam has not and does not intend to apply for the REACH Authorisation of customers' uses of

products that contain European Authorisation list (Annex XIV) substances.

It is the responsibility of our customers to check the necessity of application of REACH

Authorisation, and any other relevant authorisations, for their intended uses.

Properties

Form Liquid

Storage instructions Shipped at Room Temperature. Store at Room Temperature.

Storage buffer pH: 7.50

Constituents: 0.22% Beta glycerophosphate, 0.18% Sodium orthovanadate, 5% Sodium

deoxycholate, 0.38% EGTA, 1% Sodium lauryl sulfate, 6.1% Tris, 0.29% EDTA, 8.8% Sodium

chloride, 1.12% Sodium pyrophosphate decahydrate, 10% Nonylphenol, ethoxylated

Applications

The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab156034 in the following tested applications.

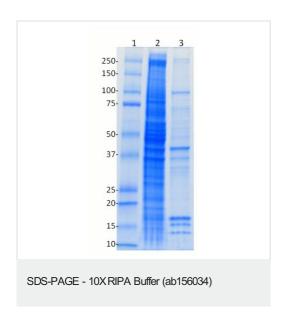
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB		Use at an assay dependent concentration. Suggested working concentration: 1X

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Application	Abreviews	Notes
ELISA		Use at an assay dependent concentration. Suggested working concentration: 1X
SDS-PAGE		Use at an assay dependent concentration. Suggested working concentration: 1X
IP		Use at an assay dependent concentration. Suggested working concentration: 1X

Images



HeLa cell extraction using ab156034.

2.5 million HeLa cells were lysed on ice for 15 minutes with 0.5 mL of 1X ab156034. Next the sample was centrifuged at 14,000 rpm at 4°C for 15 minutes: the supernatant (= cleared lysate) was removed and the pellet (= insoluble material) was resuspended in 0.5 mL lysis buffer and solubilized by sonication. Equivalent loads of the cleared lysate and solubilized pellet were analyzed by SDS-PAGE and Coomassie stain.

BCA protein concentration determination of the soluble and insoluble material indicates that a total of 1.1mg of protein was recovered and 82% was in the soluble cleared cell lysate.

Lane 1: MW marker

Lane 2: Cleared lysate

Lane 3: Non-soluble

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

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