

Native lysis Buffer ab156035

★★★★★ [1 Abreviews](#) [1 Image](#)

Overview

Product name	Native lysis Buffer
Tested applications	Suitable for: WB, Sandwich ELISA, SDS-PAGE, IP
General notes	Abcam's 1X Native lysis buffer is an efficient means of cell lysis and protein solubilization for both adherent and suspension cultured mammalian cells. This reagent extracts cytoplasmic, nuclear and membrane proteins. The detergent used in this buffer is relatively gentle and in many cases allows solubilization of protein complexes that retain enzymatic activity. On the flip side, the gentle nature of this lysis buffer means that it may result in extractions that are less complete than lysis buffers using more stringent detergents (e.g. ab156034). Native lysis buffer is compatible with many downstream applications, including enzyme activity assays, SDS-PAGE, Western blot, immunoprecipitation, ELISA and BCA assays.

Properties

Form	Liquid
Storage instructions	Store at +4°C.
Storage buffer	pH: 7.50 Constituents: 0.02% Beta glycerophosphate, 0.02% Sodium orthovanadate, 0.04% EGTA, 0.6% HEPES, 0.03% EDTA, 0.58% Sodium chloride, 1.5% Dodecyl maltoside, 0.11% Sodium pyrophosphate decahydrate

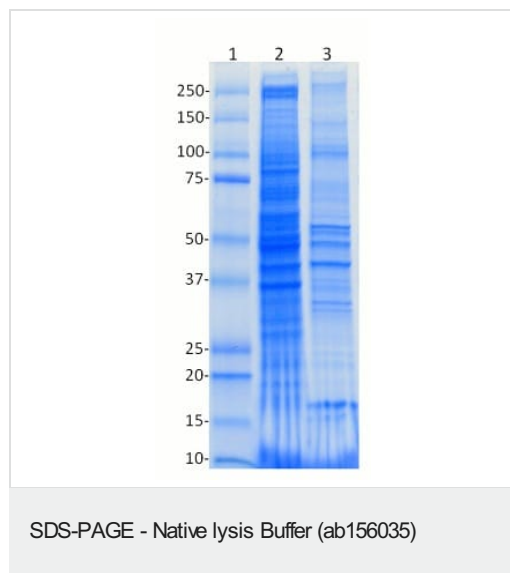
Applications

The Abpromise guarantee Our **Abpromise guarantee** covers the use of ab156035 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.
Sandwich ELISA		Use at an assay dependent concentration.
SDS-PAGE		Use at an assay dependent concentration.

Application	Abreviews	Notes
IP		Use at an assay dependent concentration.

Images



HeLa cell extraction using ab156035.

2.5 million HeLa cells were lysed on ice for 15 minutes with 0.5 mL of 1X ab156035. 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 1.15mg of protein was recovered and 70% was in the soluble cleared cell lysate.

Lane 1: MW marker

Lane 2: Cleared lysate

Lane 3: Non-soluble

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