

Product Datasheet

Ramos Cell Nuclear Extract (orb348699)



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Descriptionnts. Ramos Cell Nuclear Extract

Conjugation Unconjugated

Tested
Applications

ChIP, IP, WB

Preservatives Preservative: None. Stabilizer: None. 1X

RIPA Buffer with HALT Protease and

Phosphatase Inhibitors

Form/Appearance Liquid (sterile filtered)

Concentration 4.0 mg/ml

Storage Store vial at -70° C or COLDER. For

extended storage, aliquot Nuclear Extract

to minimize freeze/thaw cycles.

Note For research use only

Application notes Multi-purpose Ramos nuclear extracts are

especially prepared as positive control for multiple assays including western blot, immunoprecipitation (IP), capture ELISA or other assays requiring native protein sample. For separation by SDS-PAGE and subsequent western blot analysis, lysates should be diluted by user to desired concentration in SDS-PAGE buffer with 2-mercaptoethanol or dithiothreitol as the reducing agent and heated to 95° C for 5 minutes. Sample is ready for use in immunoprecipitation and ELISA experiments, conditions should be optimized by the user. Rockland recommends its TrueBlot IP reagents for

immunoprecipitation experiments.

Purity Ramos cells were grown in RPMI-1640

medium supplemented with 10% fetal bovine serum. Cells were washed with PBS and then incubated on ice in modified RIPA buffer, containing 150 mM sodium chloride, 50 mM Tris HCl, pH 7.4, 1 mM EDTA, 1.0% NP-40, 0.5% sodium deoxycholic acid , 0.1% SDS and 0.01% (w/v) sodium azide to lyse the cells. Protein integrity was ensured using a cocktail of protease inhibitors with broad specificity for the inhibition of aspartic, cysteine, and serine proteases as well as aminopeptidases (0.1 mM AEBSF HCl, 0.08 μ M Aprotinin, 5 μ M Bestatin, 1.5 μ M E-64, 2 μ M Leupeptin Hemisulfate, 1 μ M

Panatatin A). Phaanbatasa inbibitara 1 mM