

Produktinformation



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Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



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Datasheet for W10-001-GL8

C2C12 Cell Nuclear Extract

Overview

Description:	C2C12 Cell Nuclear Extract - W10-001-GL8
Item No.:	W10-001-GL8
Size:	200 μg
Origin:	Mouse

Product Details

Background:	Multi-purpose C2C12 nuclear extracts produced by Rockland Immunochemicals are derived from cell lines or tissues using highly refined extraction protocols to ensure exceptionally high quality, protein integrity and lot-to-lot reproducibility. All extracts are tested by SDS-PAGE using 4-20% gradient gels.
Synonyms:	C2C12 lysate nuclear extract, cell lysate, C2C12 Nuclear lysate
Species of Origin:	Mouse

Target Details

Purity/Specificity: C2C12 cells were grown in Dulbecco's medium supplemented with 10% fetal bovine serum.

Cells were washed with PBS and then incubated on ice in modified RIPA buffer 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 Pepstatin A). Phosphatase inhibitors 1 mM NaF and 1 mM Na3VO4 were also added. Cell debris was removed by centrifugation. Protein concentration was determined by a modified Lowry assay using a commercially available kit. Protein concentration was adjusted to 4 mg/ml in RIPA buffer containing protease and phosphatase inhibitors.

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Application Details

Application Note: Multi-purpose C2C12 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-

www.rockland.com Page 1 of 3





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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.

Assay Dilutions:	All assays should be optimized by the user. Recommended dilutions (if any) may be listed below.
ChIP:	User Optimized
IP:	User Optimized
WB:	User Optimized

Cell Line Data

Cell Line:	Mouse - Myoblast
Lysate Fractionation:	Nuclear Extract
Lysate Stimulation:	Not Stimulated
Culture Type:	Tissue Culture
Induction:	None (Control)

Formulation

Physical State:	Liquid (sterile filtered)
Concentration:	4.0 by BCA assay
Buffer:	1X RIPA Buffer with HALT Protease and Phosphatase Inhibitors
Preservative:	None
Stabilizer:	None

Shipping & Handling

Shipping Condition:	Dry Ice
Storage Condition:	Store vial at -70° C or COLDER. For extended storage, aliquot Nuclear Extract to minimize freeze/thaw cycles.
Expiration:	Expiration date is three (3) months from date of receipt.

Disclaimer

www.rockland.com Page 2 of 3





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www.rockland.com Page 3 of 3