

# Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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# Lieferung & Zahlungsart

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- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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#### Datasheet for W12-001-GM1

## PC-12 Cell Nuclear Extract

#### **Overview**

Description:	PC-12 Cell Nuclear Extract - W12-001-GM1
Item No.:	W12-001-GM1
Size:	200 μg
Origin:	Rat

#### **Product Details**

Background:	Multi-purpose PC-12 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:	PC-12 lysate nuclear extract, cell lysate, PC12 Nuclear lysate
Species of Origin:	Rat

#### **Target Details**

**Purity/Specificity:** PC-12 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 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  $\mu$ M Aprotinin, 5  $\mu$ M Bestatin, 1.5  $\mu$ M E-64, 2  $\mu$ M Leupeptin Hemisulfate, 1  $\mu$ 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.

### **Application Details**

**Application Note:** Multi-purpose PC-12 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-

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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:	Rat - adrenal gland
Lysate Fractionation:	Nuclear Extract
Lysate Stimulation:	Not Stimulated
Culture Type:	Tissue Culture
Induction:	None (Control)

# **Formulation**

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

# **Shipping & Handling**

<b>Shipping Condition:</b>	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**

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