

Datasheet for W09-001-370

Jurkat Whole Cell Lysate

Overview

Description:	Jurkat Whole Cell Lysate - W09-001-370
Item No.:	W09-001-370
Size:	500 µg
Applications:	SDS-PAGE, WB
Origin:	Human

Product Details

Background:	Ready-to-use whole cell lysates 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 and immunoblot analysis using antibodies to key cell signaling components to confirm the presence of both high molecular weight and low molecular weight proteins.
Synonyms:	Jurkat Whole Cell Lysate, Jurkat WCL, Jurkat Lysate
Species of Origin:	Human

Target Details

Purity/Specificity:	The cells were grown in RPMI 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 µM Aprotinin, 5 µM Bestatin, 1.5 µM E-64, 2 µM Leupeptin Hemisulfate, 1 µM 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 2 mg/ml and then an equal volume of 2X SDS-PAGE sample buffer was added.
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Application Details

Tested Applications:	SDS-PAGE, WB
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Application Note: W09-001-370 has been tested by SDS-PAGE and western blot. Ready-to-use lysates are especially prepared as positive controls for separation by SDS-PAGE and subsequent western blot analysis. Lysates are prepared in denaturing buffer WITHOUT dissociating agents (i.e. no 2-mercaptoethanol or dithiothreitol has been added). Heat lysate to 95° C for 5 minutes and rapidly cool. If dissociating conditions are desired, add reducing agent prior to heating. The recommended loading volume per lane is 10-20 µl depending on the size format of your gel.

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

WB: User Optimized

Cell Line Data

Cell Line: Human T Lymphocyte (Acute T Leukemia)

Lysate Fractionation: Whole Cell Lysate

Lysate Stimulation: Not Stimulated

Culture Type: Tissue Culture

Induction: None (Control)

Formulation

Physical State: Liquid (sterile filtered)

Concentration: 1.0 mg/ml by BCA assay

Buffer: 1X SDS-PAGE Sample Buffer (62.5 mM Tris HCl, 2% SDS, 10% Glycerol and 0.005% bromophenol blue, pH 6.8)

Preservative: None

Stabilizer: 10% (v/v) Glycerol

Shipping & Handling

Shipping Condition: Dry Ice

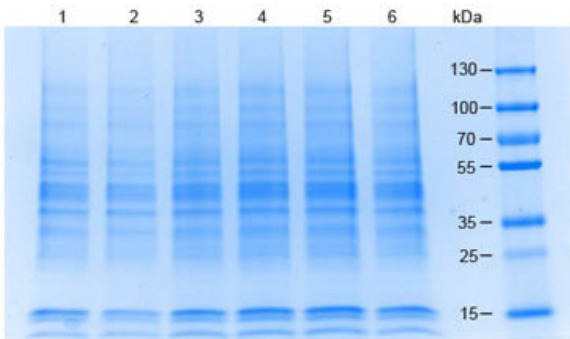
Storage Condition: Store vial at -70° C or COLDER. For extended storage, aliquot contents to minimize freeze/thaw cycles.

Expiration: Expiration date is three (3) months from date of receipt.

Images

SDS-PAGE

Coomassie stained SDS-PAGE of 20 µg of (1) HeLa WCL, (2) Jurkat WCL, (3) HEK 293 WCL, (4) MCF-7 WCL, (5) A 549 WCL, (6) Raji WCL separated using a 4-20% gradient gel under reducing conditions. Molecular weight standards are shown.

**Disclaimer**

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