

## Datasheet for W09-000-363

## Hep2 Whole Cell Lysate

### Overview

<b>Description:</b>	Hep2 Whole Cell Lysate - W09-000-363
<b>Item No.:</b>	W09-000-363
<b>Size:</b>	500 µg
<b>Applications:</b>	SDS-PAGE
<b>Origin:</b>	Human

### Product Details

<b>Background:</b>	Hep2 Whole Cell Lysate 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.
<b>Synonyms:</b>	Hep2, Lysate, Whole Cell Lysate, Hep2 Lysate
<b>Species of Origin:</b>	Human

### Target Details

<b>Purity/Specificity:</b>	Hep2 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 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

<b>Tested Applications:</b>	SDS-PAGE
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<b>Application Note:</b>	W09-000-363 has been tested by SDS-PAGE. 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.
<b>Assay Dilutions:</b>	All assays should be optimized by the user. Recommended dilutions (if any) may be listed below.
<b>WB:</b>	User Optimized

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## Cell Line Data

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<b>Cell Line:</b>	Human HEp2 (HeLa contaminant)
<b>Lysate Fractionation:</b>	Whole Cell Lysate
<b>Lysate Stimulation:</b>	Not Stimulated
<b>Culture Type:</b>	Tissue Culture
<b>Induction:</b>	None (Control)

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## Formulation

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<b>Physical State:</b>	Liquid
<b>Concentration:</b>	1.0 mg/mL by modified Lowry assay
<b>Buffer:</b>	1X SDS-PAGE Sample Buffer (62.5 mM Tris HCl, 2% SDS, 10% Glycerol and 0.005% bromophenol blue, pH 6.8)
<b>Preservative:</b>	None
<b>Stabilizer:</b>	10% (v/v) Glycerol

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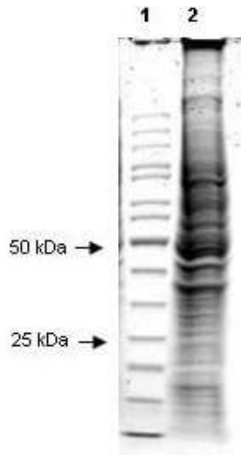
## Shipping & Handling

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

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## Images

**SDS-PAGE**

Coomassie stained SDS-PAGE of 20  $\mu$ l of Human Derived Hep2 Whole Cell Lysate (Ready-to-Use) separated in a 4-20% gradient gel under reducing conditions (lane 2). Molecular weight standards are shown in lane 1.

**Disclaimer**

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