

**Datasheet for W09-001-GJ5**

## Hep-G2 Lysate

### Overview

<b>Description:</b>	Hep-G2 - Whole Cell Lysate - W09-001-GJ5
<b>Item No.:</b>	W09-001-GJ5
<b>Size:</b>	500 µg
<b>Applications:</b>	SDS-PAGE, WB, Cellular Assay
<b>Origin:</b>	Human

### Product Details

<b>Background:</b>	Multi-purpose Hep-G2 Whole Cell Lysates produced by Rockland Immunochemicals are derived from cell lines 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.
<b>Synonyms:</b>	HepG2 Lysate, Cell Lysate, Hep-G2 Lysate
<b>Species of Origin:</b>	Human
<b>Clone ID:</b>	Hep-G2

### Target Details

<b>Purity/Specificity:</b>	Hep-G2 cells were grown in Eagle's Minimum Essential 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 include sodium fluoride, sodium orthovanadate, sodium pyrophosphate and β-glycerophosphate. 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 with 1X RIPA buffer including protease and phosphatase inhibitors.
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### Application Details

<b>Tested Applications:</b>	SDS-PAGE, WB
<b>Suggested Applications:</b>	Cellular Assay (Based on references)
<b>Application Note:</b>	W09-001-GJ5 has been tested by SDS-PAGE and western blot. Multi-purpose Hep-G2 Whole Cell Lysate is 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.
<b>Assay Dilutions:</b>	All assays should be optimized by the user. Recommended dilutions (if any) may be listed below.
<b>ChIP:</b>	User Optimized
<b>IP:</b>	User Optimized
<b>WB:</b>	User Optimized
<b>Other:</b>	User Optimized

## Cell Line Data

<b>Cell Line:</b>	Human - hepatocellular carcinoma
<b>Lysate Fractionation:</b>	Whole Cell Lysate
<b>Lysate Stimulation:</b>	Not Stimulated
<b>Culture Type:</b>	Tissue Culture
<b>Induction:</b>	None (Control)

## Formulation

<b>Physical State:</b>	Liquid (sterile filtered)
<b>Concentration:</b>	1.0 mg/mL by BCA assay
<b>Buffer:</b>	1X RIPA Buffer with HALT Protease and Phosphatase Inhibitors
<b>Preservative:</b>	None
<b>Stabilizer:</b>	None

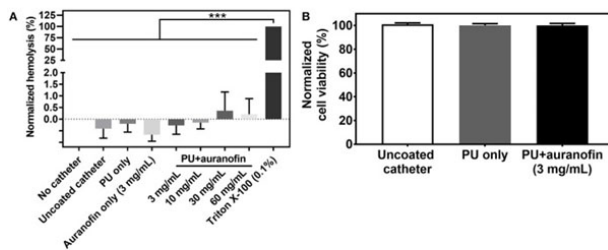
## Shipping & Handling

<b>Shipping Condition:</b>	Dry Ice
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**Storage Condition:** Store Hep-G2 Whole Cell Lysate at -70° C or COLDER. For extended storage, whole cell lysate to minimize freeze/thaw cycles.

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

## Images



### Figure

Cytotoxicity of PU+auranofin coatings. (A) Percent normalized hemolysis of hRBCs exposed to uncoated catheters, PU only, auranofin only (formulated at a 3 mg/mL auranofin coating concentration), PU+auranofin (formulated at 3, 10, 30, and 60 mg/mL auranofin coating concentrations) compared to negative controls of untreated hRBCs and a Triton X-100 incubated positive control. (B) Normalized HepG2 liver cell viability upon exposure to media incubated with uncoated catheters, PU only, and PU+auranofin (formulated at a 3 mg/mL auranofin coating concentration) catheters for 24 h. Data are shown as mean  $\pm$  standard deviation. Statistical significance was evaluated using one-way ANOVA ( $n = 3$ ) and is shown as \*\*\* $p < 0.001$  indicating statistical significance between the positive control (Triton X-100 with hRBCs) and other conditions tested. No statistical significance was noted between the other hemolysis conditions tested or between the different HepG2 viability conditions examined ( $p > 0.5$ ). HepG2 (p/n W09-001-GJ5). Figure 8. PMID: 30873389.

## References

- Liu, H et al. Auranofin Releasing Antibacterial and Antibiofilm Polyurethane Intravascular Catheter Coatings. *Frontiers in Cellular and Infection Microbiology* (2019)

## Disclaimer

No test method can provide total assurance that the hepatitis B virus, hepatitis C virus, human immunodeficiency virus, or any other infectious agents are absent. Thus, all blood products, including purified proteins derived from human blood sources, should be handled at Biosafety Level 2 as recommended by the CDC\NIH manual entitled Biosafety in Microbiological and Biomedical Laboratories for potentially infectious human serum, blood specimens or proteins derived from same. Source material for the human blood product supplied to your facility has been tested for the detection of HIV antibody, Hepatitis B surface antigen, antibody to Hepatitis C, HIV 1 antigen(s), antibody to HTLV - I/II, and syphilis by FDA guidelines. All units were found to be non-reactive/negative for these tests. All human blood source material is collected in FDA licensed centers and is tested with FDA approved test kits.

This product is for research use only and is not intended for therapeutic or diagnostic applications. Please contact a technical service representative for more information. All products of animal origin manufactured by Rockland Immunochemicals are derived from starting materials of North American origin. Collection was performed in United States Department of Agriculture (USDA) inspected facilities and all materials have been inspected and certified to be free of disease and suitable for exportation. All properties listed are typical characteristics and are not specifications. All suggestions and data are offered in good faith but without guarantee as conditions and methods of use of our products are beyond our control. All claims must be made within 30 days following the date of delivery. The prospective user must determine the suitability of our materials before adopting them on a commercial scale. Suggested uses of our products are not recommendations to use our products in violation of any patent or as a license under any patent of Rockland Immunochemicals, Inc. If you require a commercial license to use this material and do not have one, then return this material, unopened to: Rockland Inc., P.O. BOX 5199, Limerick, Pennsylvania, USA.