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Lieferung & Zahlungsart

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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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Datasheet for W07-000-359

CHO/K1 Whole Cell Lysate**Overview**

| | |
|----------------------|--|
| Description: | CHO/K1 Whole Cell Lysate - W07-000-359 |
| Item No.: | W07-000-359 |
| Size: | 500 µg |
| Applications: | SDS-PAGE, WB |
| Origin: | Chinese Hamster |

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: | CHO/K1 Whole Cell Lysate, CHO K1 WCL, CHO-K1 lysate, CHOK1 lysate |
| Species of Origin: | Chinese Hamster |

Target Details

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|----------------------------|---|
| Purity/Specificity: | The cells were grown in Ham's F12 medium containing 2 mM L-Gln and 1.5 g/L sodium bicarbonate supplemented with 10% FBS (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. |
|----------------------------|---|

Application Details

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|-----------------------------|--------------|
| Tested Applications: | SDS-PAGE, WB |
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| Application Note: | W07-000-359 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

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|------------------------------|-----------------------------|
| Cell Line: | Chinese Hamster Ovary (CHO) |
| Lysate Fractionation: | Whole Cell Lysate |
| Lysate Stimulation: | Not Stimulated |
| Culture Type: | Tissue Culture |
| Induction: | None (Control) |

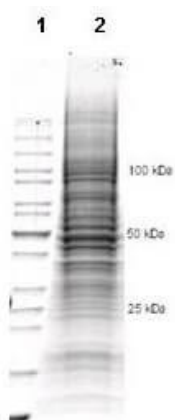
Formulation

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|------------------------|--|
| Physical State: | Liquid |
| 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

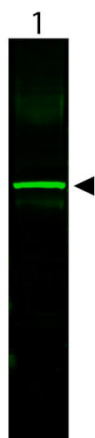
| | |
|----------------------------|--|
| Shipping Condition: | Dry Ice |
| Storage Condition: | Store vial at -70° C or COLDER. For extended storage, aliquot contents to minimize freeze/thaw cycles. |
| Expiration: | Expiration date is three (3) months from date of receipt. |

Images



SDS-PAGE

Coomassie stained SDS-PAGE of 35 μ l of CHO 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.



Western Blot

Western Blot showing detection of alpha tubulin in lane 1. CHO/K1 Whole Cell Lysate (10 μ g) (p/n W07-000-359) was run on a 4-20% gel, then transferred to 0.45 μ m nitrocellulose. After blocking with 1% BSA-TTBS (p/n MB-013, diluted to 1X) 30 min at 20°C, primary antibody was used at 1:1000 overnight at 4°C. Anti-Rabbit IgG (H&L) (GOAT) antibody IRDye800CW® (p/n 611-131-002) secondary antibody was used at 1:20,000 in Blocking Buffer for Fluorescent Western Blotting (p/n MB-070) and imaged on the LiCor Odyssey imaging system. Arrow indicates correct 50 kDa molecular weight position expected for alpha tubulin.

Disclaimer

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.