

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten! See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com



Datasheet for 610-4302

Mouse IgG (H&L) Secondary Antibody Peroxidase Conjugated

Overview

Description:	Rabbit Anti-Mouse IgG (H&L) Antibody Peroxidase Conjugated - 610-4302
Item No.:	610-4302
Size:	2 mg
Applications:	ELISA, WB, IF, Other
Reactivity:	Mouse
Host Species:	Rabbit

Product Details

п	_	_	١.	_	 _	u	 _1	
$\boldsymbol{-}$	ч	•	10	5	v	u	 u	•

Secondary antibodies bind to the primary antibody to assist in detection, sorting and purification of target antigens. To enable detection, the secondary antibody must have specificity for the antibody species and isotype of the primary antibody being used and generally is conjugated. Rockland produces highly active antibodies and conjugates against mouse immunoglobulins. Anti-Mouse Secondary Antibodies are affinity-purified polyclonal antibodies with well-characterized specificity for mouse immunoglobulin classes, subclasses, and fragments. They are available as unlabeled antibodies as well as conjugates of alkaline phosphatase (AP), horseradish peroxidase (HRP), fluorescent conjugates, and biotin. This Anti-MOUSE IgG generated in rabbit is designed to detect heavy and light chains (H&L) and has been Peroxidase (HRP) conjugated. Rigorous quality control testing ensures that the finished product meets or exceeds out high standards for optimum performance in your assays. These secondary antibodies are used throughout various types of assays, including ELISA or Western Blot, Immunohistochemistry, Flow Cytometry. Optimal secondary antibody requires knowledge of the detection assay.

Synonyms:

Rabbit Anti-Mouse IgG Secondary Antibody Peroxidase Conjugated, Rabbit Anti-Mouse IgG Secondary Antibody HRP Conjugated, RAM-HRP, Anti-mouse secondary antibody, anti-mouse HRP antibody, horseradish peroxidase conjugated secondary antibody, anti-mouse horseradish peroxidase conjugated secondary antibody

Host Species:	Rabbit
Specificity:	IgG (H&L)
Conjugate:	Peroxidase (HRP)
Clonality:	Polyclonal

www.rockland.com Page 1 of 5



Format: IgG

Target Details

Reactivity:	Mouse
Immunogen:	Mouse IgG whole molecule
Purity/Specificity:	HRP Secondary Antibody Conjugate was prepared from monospecific antiserum by immunoaffinity chromatography using Mouse IgG coupled to agarose beads. Assay by immunoelectrophoresis resulted in a single precipitin arc against anti-Peroxidase, anti-Rabbit Serum, Mouse IgG and Mouse Serum.

Application Details

Tested Applications:	ELISA, WB
Suggested Applications:	IF, Other (Based on references)
Application Note:	Anti-Mouse secondary antibody conjugated to horseradish peroxidase (HRP) generated in rabbit detects specifically Mouse IgG whole molecule. Anti-Mouse IgG peroxidase antibody has been tested by ELISA and western blot and is suitable for ELISA, Sandwich ELISA, titration assays, western-blot, immunoprecipitation, Immunohistochemistry as well as other HRP antibody based assays. Specific conditions for reactivity and signal detection should be optimized by the end user.
Assay Dilutions:	All assays should be optimized by the user. Recommended dilutions (if any) may be listed below.
ELISA:	1:50,000 - 1:300,000
IHC:	1:500 - 1:2,500
WB:	1:10,000 - 1:30,000

Formulation

Physical State:	Lyophilized
Concentration:	2.0 mg/mL by UV absorbance at 280 nm
Buffer:	0.02 M Potassium Phosphate, 0.15 M Sodium Chloride, pH 7.2
Preservative:	0.01% (w/v) Gentamicin Sulfate. Do NOT add Sodium Azide!
Stabilizer:	10 mg/mL Bovine Serum Albumin (BSA) - Immunoglobulin and Protease free
Reconstitution Volume:	1.0 mL

www.rockland.com Page 2 of 5

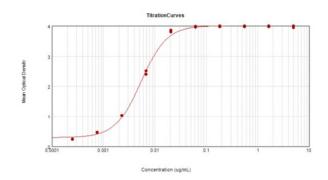


Reconstitution Buffer: Restore with deionized water (or equivalent)

Shipping & Handling

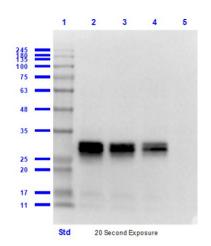
Shipping Condition:	Ambient
Storage Condition:	Store mouse secondary antibody conjugate at 4° C prior to restoration. For extended storage aliquot antibody and freeze at -20° C or below. Avoid cycles of freezing and thawing. Centrifuge product if not completely clear after standing at room temperature. This product is stable for several weeks at 4° C as an undiluted liquid. Dilute only prior to immediate use.
Expiration:	Expiration date is one (1) year from date of receipt.

Images



ELISA

ELISA Results of Rabbit Anti-Mouse IgG Peroxidase Antibody tested against purified Mouse IgG protein. Each well was coated in duplicate with 10 μ g of Mouse IgG (p/n 010-0102). The starting dilution of antibody was 5μ g/ml and the X-axis represents the Log10 of a 3-fold dilution. The titer is 1:185,000. This titration is a 4-parameter curve fit where the IC50 is defined as the titer of the antibody. Assay performed using 3% fish gelatin as blocking buffer and TMB substrate p/n TMBE-1000.

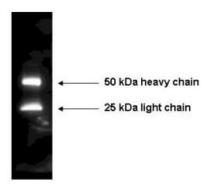


Western Blot

Western blot using Rabbit Anti-Mouse IgG Peroxidase Conjugated Antibody and Mouse Anti-GFP Antibody. Lane 1: Opal Prestained Molecular Weight Marker (p/n MB-210-0500). Lane 2: HeLa WC Lysate+GFP protein (p/n W09-000-364 [10µg]/ p/n 000-001-215 [0.1µg]). Lane 3: HeLa WC Lysate+GFP protein [10µg/0.066µg]. Lane 4: HeLa WC Lysate+GFP protein [10µg/0.033µg]. Lane 5: HeLa Whole Cell Lysate (p/n W09-000-364) [10µg]. Primary Antibody: Anti-GFP at 1:1000 overnight at 2-8°C. Secondary Antibody: Rabbit Anti-Mouse IgG HRP (p/n 610-4302) at 1:40,000 for 30mins at RT. Block: BlockOut Buffer (p/n MB-073) for 60mins at RT. Expected MW: ~27kDa.

www.rockland.com Page 3 of 5





Western Blot

Western Blot of Rockland's Anti-Mouse IgG Antibody Peroxidase Conjugated detecting mouse IgG. Mouse IgG (1.0 mg) was separated on a 4-20% gradient gel and then transferred to Whatman Protran BA 85 nitrocellulose (0.45 µm pore size), and blocked with 5% BLOTTO (p/n B501-0500) o/n at 4° C. The membrane was then probed with a 1:2,500 dilution of HRP Rb-a-Mouse IgG [H&L] (p/n 610-4302) for 45 min at RT. FemtoMax Super Sensitive Chemiluminescent HRP Substrate (p/n FEMTOMAX-110) was added and the image was captured using a BioSpectrum Imaging System with a BioChemi 500 -28C cooled 4.0 Megapixel CCD Camera from UVP (www.uvp.com, Upland, CA). Binning: 5X5. Exposure time: 2 min. Aperture open. No emission or excitation filters used.

References

- van Soest DMK et al. Mitochondrial H2O2 release does not directly cause damage to chromosomal DNA. Nat Commun.
 (2024)
- Beitari S et al. Effectiveness of VSV vectored SARS-CoV-2 spike when administered through intranasal, intramuscular or a combination of both. *Sci Rep.* (2023)
- Srivastava AK et al. Ferritin microheterogeneity, subunit composition, functional, and physiological implications. *Sci Rep.* (2023)
- Amanat F et al. Immunity to Seasonal Coronavirus Spike Proteins Does Not Protect from SARS-CoV-2 Challenge in a Mouse Model but Has No Detrimental Effect on Protection Mediated by COVID-19 mRNA Vaccination. J Virol. (2023)
- González MF et al. Extracellular vesicles from gastric epithelial GES-1 cells infected with Helicobacter pylori promote changes in recipient cells associated with malignancy. *Front Oncol.* (2022)
- Lim JH et al. Palmitoyl@RGD promotes the expression of dermal@epidermal junction components in HaCaT cells. *Mol Med Rep.* (2022)
- Amanat F et al. The plasmablast response to SARS-CoV-2 mRNA vaccination is dominated by non-neutralizing antibodies that target both the NTD and the RBD. *medRxiv*. (2021)
- Amanat F et al. Introduction of Two Prolines and Removal of the Polybasic Cleavage Site Lead to Higher Efficacy of a Recombinant Spike-Based SARS-CoV-2 Vaccine in the Mouse Model. *mBio.* (2021)
- Amanat F et al. SARS-CoV-2 mRNA vaccination induces functionally diverse antibodies to NTD, RBD, and S2. Cell. (2021)
- Amanat F et al. Vaccination with SARS-CoV-2 variants of concern protects mice from challenge with wild-type virus. PLoS
 One. (2021)
- Wang TS et al. Endolysosomal Targeting of Mitochondria Is Integral to BAX-Mediated Mitochondrial Permeabilization during Apoptosis Signaling. *Dev Cell.* (2020)

www.rockland.com Page 4 of 5



- Amanat F et al. An in vitro microneutralization assay for SARS-CoV-2 serology and drug screening. Curr Protoc Microbiol.
 (2020)
- Amanat F et al. Introduction of two prolines and removal of the polybasic cleavage site leads to optimal efficacy of a recombinant spike based SARS-CoV-2 vaccine in the mouse model. ASM (2020)
- Jiang C et al. Regulation of Mitochondrial Respiratory Chain Complex Levels, Organization, and Function by Arginyltransferase 1. Front Cell Dev Biol (2020)
- Wakabayashi R et al. Transcutaneous codelivery of tumor antigen and resiquimod in solid-in-oil nanodispersions promotes antitumor immunity. ACS Biomater Sci Eng. (2019)
- Fegan, JE et al. Utility of Hybrid Transferrin Binding Protein Antigens for Protection Against Pathogenic Neisseria Species. Frontiers in Immunology (2019)
- Kirkpatrick et al. The influenza virus hemagglutinin head evolves faster than the stalk domain. Scientific Reports (2018)
- Platt TL et al. Obesity, diabetes, and leptin resistance promote tau pathology in a mouse model of disease. *Neuroscience* (2016)
- Beckett TL et al. Postmortem Pittsburgh Compound B (PiB) binding increases with Alzheimer's disease progression. Journal of Alzheimer's Disease: Jad (2012)
- O'shannessy, DJ et al. Characterization of the human folate receptor alpha via novel antibody-based probes. *Oncotarget* (2011)
- Beckett TL et al. Effects of nonsteroidal anti-inflammatory drugs on amyloid-beta pathology in mouse skeletal muscle. Neurobiology of Disease (2010)
- McCall LI et al. Localization and induction of the A2 virulence factor in Leishmania: evidence that A2 is a stress response protein. Mol Microbiol. (2010)

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.

www.rockland.com Page 5 of 5