

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 <u>mail@szabo-scandic.com</u> www.szabo-scandic.com



Datasheet for 18-8814-31 Goat TrueBlot[®]: Anti-Goat IgG HRP

Overview

Description:	Goat TrueBlot®: Anti-Goat IgG HRP - 18-8814-31
Item No.:	18-8814-31
Size:	50 μL
Applications:	IP, WB
Reactivity:	Goat
Host Species:	Mouse

Product Details

Background:	Goat IgG TrueBlot [®] is a unique horseradish peroxidase conjugated Anti-Goat IgG monoclonal secondary antibody. Goat IgG TrueBlot [®] enables detection of immunoblotted target protein bands, without hindrance by interfering immunoprecipitating immunoglobulin heavy and light chains. It is easy to generate publication-quality IP/Western Blot data with Goat IgG TrueBlot [®] , simply substitute the conventional HRP Anti-Goat IgG blotting reagent with Goat IgG TrueBlot [®] and follow the prescribed protocol for sample preparation and immunoblotting. Goat IgG TrueBlot [®] is ideal for use in protocols involving immunoblotting of immunoprecipitated proteins. TrueBlot preferentially detects the non-reduced form of Goat IgG over the reduced, SDS-denatured form of IgG. When the immunoprecipitate is fully reduced immediately prior to SDS-gel electrophoresis, reactivity of Goat IgG TrueBlot [®] with the 55 kDa heavy chains and the 23 kDa light chains of the immunoprecipitating antibody is minimized thereby eliminating interference by the heavy and light chains of the immunoprecipitating antibody in IP/immunoblot applications. Applications include studies examining post-translational modification (e.g., phosphorylation or acetylation) or protein-protein interactions.
Synonyms:	Anti-Goat IgG HRP, TrueBlot, HRP TrueBlot ULTRA, Peroxidase TrueBlot, TrueBlot for IP/WB, TrueBlot for immunoprecipitation, TrueBlot for western blotting
Host Species:	Mouse
Conjugate:	Peroxidase (HRP) ULTRA
Clonality:	Monoclonal
Clone ID:	eB270
Format:	IgG
Detection Kit Type:	Immunoprecipitation Kit



3.1

www.rockland.com tech@rockland.com +1 484.791.3823

F/P Ratio:

Target Details

Reactivity:	Goat
Purity/Specificity:	Goat TrueBlot [®] Antibody Peroxidase Conjugate was prepared from tissue culture supernatant by Protein G affinity chromatography. Assay by Immunoelectrophoresis resulted in a single precipitin arc against Anti-Goat Serum. Reactivity is observed against native Goat IgG by both Western blot and ELISA.
Relevant Links:	 TrueBlot HRP Product Protocols TrueBlot IP Set Protocol

Application Details

be used for detection in immunoblotting assays that do not employ immunoprecipitation. Go IgG TrueBlot® is provided as 1000X solution. To conserve reagent, we recommend incubating the blots from minigels in sealed bags (removing as much air as possible) with minimal volum (2-3 mLs). If used conservatively at 2.5mls/blot will yield enough reagent for 20 blots. All recommended dilutions for listed applications are intended as an initial recommendation, specific conditions for each protein and antibody combination should be specifically optimize by the end user. Note that there are three key procedural considerations: 1. Protein A or G beads may be used with the Mouse, Goat and Sheep TrueBlot secondaries, but not with the	Tested Applications:	IP, WB
listed below.	Application Note:	recommended dilutions for listed applications are intended as an initial recommendation, specific conditions for each protein and antibody combination should be specifically optimized by the end user. Note that there are three key procedural considerations: 1. Protein A or G beads may be used with the Mouse, Goat and Sheep TrueBlot secondaries, but not with the Rabbit TrueBlot secondary. Use of protein A or G beads with the Rabbit TrueBlot secondary. 2. Immunoprecipitate should be completely reduced. 3. BLOTTO/Milk
WB: 1:1000	Assay Dilutions:	
	WB:	1:1000

Formulation

Physical State:	Liquid (sterile filtered)
Concentration:	1 mg/mL by UV absorbance at 280 nm
Buffer:	0.02 M Potassium Phosphate, 0.15 M Sodium Chloride, pH 7.2
Preservative:	Proclin is added as an antimicrobial agent.



Stabilizer: 0.1 mg/ml Bovine Serum Albumin (BSA) - IgG and Protease free, 50% (v/v) Glycerol

Shipping & Handling

Shipping Condition:	Wet Ice
Storage Condition:	Store at -20°C. This product is guaranteed for 6 months upon receipt, when handled and stored as instructed.
Expiration:	Expiration date is six (6) months from date of receipt.

Images



Western Blot

Goat TrueBlot[®] IP / Western Blot: Jurkat cell lysate (0.5 ml of 1x10e7 cells/ml) was incubated with goat anti-human Stat1 and immunoprecipitated using Protein G. Precipitate from 5x10e5 cells was subjected to electrophoresis, transferred to a PVDF membrane, and Western blotted with anti-Stat1 using Goat TrueBlot[®]: Anti-Goat IgG HRP (lane 1) and conventional HRP-conjugated anti-goat polyclonal antibody (lane 2).

References

- Jones VT et al. Inhibition of autocrine HGF maturation overcomes cetuximab resistance in colorectal cancer. Cell Mol Life Sci. (2024)
- Wu SA et al. The mechanisms to dispose of misfolded proteins in the endoplasmic reticulum of adipocytes. *Nat Commun.* (2023)
- Vignogna RC et al. Evolutionary rescue of phosphomannomutase deficiency in yeast models of human disease. *Elife*. (2022)
- Malgorzata Bodaszewska-Lubas et al. Dominant-Negative Form of SIGIRR: SIGIRR∆E8 Promotes Tumor Growth Through Regulation of Metabolic Pathways. J Interferon Cytokine Res. (2022)
- Chan ES et al. Allosteric potentiation of GABAA receptor single-channel conductance by netrin-1 during neuronalexcitation-induced inhibitory synaptic homeostasis. *Cell Rep.* (2022)



- Ali SR et al. Angiocrine IGFBP3 spatially coordinates IGF signaling during neonatal cardiac regeneration. *bioRxiv Preprint* (2021)
- Englert H et al. Defective NET clearance contributes to sustained FXII activation in COVID-19-associated pulmonary thrombo-inflammation. *eBioMedicine* (2021)
- Ali S et al. Angiocrine IGFBP3 spatially coordinates IGF signaling during neonatal cardiac regeneration. *bioRxiv Preprint* (2021)
- Ochiai K et al. Protocol for in vitro BCR-mediated plasma cell differentiation and purification of chromatin-associated proteins. *STAR Protoc.* (2021)
- Kubo Y et al. Periostin and tenascin-C interaction promotes angiogenesis in ischemic proliferative retinopathy. *Sci Rep.* (2020)
- Jäger MA, De La Torre C, Arnold C, et al. Assembly of vascular smooth muscle cells in 3D aggregates provokes cellular quiescence. *Exp Cell Res.* (2020)
- Boonying W, Joselin A, Huang E, et al. Pink1 regulates FKBP5 interaction with AKT/PHLPP and protects neurons from neurotoxin stress induced by MPP. *J Neurochem*. (2019)
- Graves-Deal et al. Broad-spectrum receptor tyrosine kinase inhibitors overcome de novo and acquired modes of resistance to EGFR-targeted therapies in colorectal cancer. *Oncotarget* (2019)
- Okumura T, Ohuchida K, Kibe S, et al. Adipose tissue-derived stromal cells are sources of cancer-associated fibroblasts and enhance tumor progression by dense collagen matrix. *Int J Cancer*. (2019)
- Cayrol C et al. Environmental allergens induce allergic inflammation through proteolytic maturation of IL-33 *Nat Immunol* (2018)
- Clark et al. Selected missense mutations impair frataxin processing in Friedreich ataxia. Annals of Clinical and Translational Neurology (2017)
- Liu et al. Improvement of Pharmacokinetic Profile of TRAIL via Trimer-Tag Enhances its Antitumor Activity in vivo. *Scientific Reports* (2017)
- Garcia et al. Borrelia burgdorferi BBK32 Inhibits the Classical Pathway by Blocking Activation of the C1 Complement Complex. *PLOS Pathogens* (2016)
- Puglisi R, Maccari I, Pipolo S, Mangia F, Boitani C. The nuclear form of glutathione peroxidase 4 colocalizes and directly interacts with protamines in the nuclear matrix during mouse sperm chromatin assembly. *Spermatogenesis*. (2014)
- Miyairi I, Ziebarth J, Laxton JD, et al. Host genetics and Chlamydia disease: prediction and validation of disease severity mechanisms. *PLoS One.* (2012)
- Nelersa CM et al. High-content analysis of proapoptotic EphA4 dependence receptor functions using small-molecule libraries. J Biolmol Screen (2012)
- Medici D, Shore EM, Lounev VY, Kaplan FS, Kalluri R, Olsen BR. Conversion of vascular endothelial cells into multipotent stem-like cells. *Nat Med.* (2011)
- Vuletic S, Dong W, Wolfbauer G, Tang C, Albers JJ. PLTP regulates STAT3 and NFκB in differentiated THP1 cells and human monocyte-derived macrophages. *Biochim Biophys Acta*. (2011)



• Miyairi I, Tatireddigari VR, Mahdi OS, et al. The p47 GTPases ligp2 and lrgb10 regulate innate immunity and inflammation to murine Chlamydia psittaci infection. *J Immunol.* (2007)

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.