ANTIBODIES ONLINE

Datasheet for ABIN102181 Sheep anti-Rat IgG (Heavy & Light Chain) Antibody (TRITC) -Preadsorbed



Image

Overview

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Quantity:	2 mg
Target:	IgG
Binding Specificity:	Heavy & Light Chain
Reactivity:	Rat
Host:	Sheep
Clonality:	Polyclonal
Conjugate:	TRITC
Application:	Flow Cytometry (FACS), FLISA, Fluorescence Microscopy (FM)

Product Details

Immunogen:	Immunogen: Rat IgG whole molecule
lsotype:	lgG
Specificity:	IgG (H&L)
Characteristics:	Concentration Definition: by UV absorbance at 280 nm
Purification:	Preadsorption: Solid phase absorption
Labeling Ratio:	3.0
Target Details	
Target:	lgG
Abstract:	IgG Products

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Target Details	
Target Type:	Antibody
Background:	Synonyms: Sheep Anti-Rat IgG rhodamine Conjugated Antibody, Sheep Anti-Rat IgG Antibody TRITC Conjugation
	Background: Anti-Rat IgG (H&L) Rhodamine Antibody generated in sheep detects reactivity to
	Rat IgG. Secreted as part of the adaptive immune response by plasma B cells, immunoglobulin
	G constitutes 75 % of serum immunoglobulins. Immunoglobulin G binds to viruses, bacteria, as
	well as fungi and facilitates their destruction or neutralization via agglutination (and thereby
	immobilizing them), activation of the compliment cascade, and opsinization for phagocytosis.
	The whole IgG molecule possesses both the $F(c)$ region, recognized by high-affinity Fc receptor
	proteins, as well as the F(ab) region possessing the epitope-recognition site. Both the Heavy
	and Light chains of the antibody molecule are present. Secondary Antibodies are available in a
	variety of formats and conjugate types. When choosing a secondary antibody product,
	consideration must be given to species and immunoglobulin specificity, conjugate type,
	fragment and chain specificity, level of cross-reactivity, and host-species source and fragment
	composition. This Anti-Rat IgG is conjugated to Rhodamine.

Application Details

	IF Microscopy Dilution: 1:1,000 - 1:5,000
	Flow Cytometry Dilution: 1:500 - 1:2,500
	FLISA Dilution: 1:10,000 - 1:50,000
	multiplex analysis, including multicolor imaging, utilizing various commercial platforms.
	based plate assays (FLISA) and fluorescent western blotting. This product is also suitable for
Application Notes:	Application Note: This product is designed for immunofluorescence microscopy, fluorescence

Handling

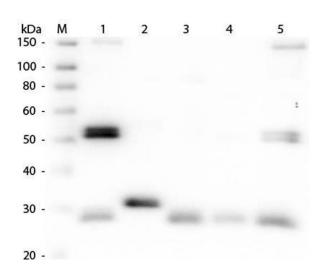
Format:	Lyophilized
Reconstitution:	Reconstitution Volume: 1.0 mL Reconstitution Buffer: Restore with deionized water (or equivalent)
Concentration:	2.0 mg/mL
Buffer:	Buffer: 0.02 M Potassium Phosphate, 0.15 M Sodium Chloride, pH 7.2 Stabilizer: 10 mg/mL Bovine Serum Albumin (BSA) - Immunoglobulin and Protease free Preservative: 0.01 % (w/v) Sodium Azide

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Handling

Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Handling Advice:	Product is photosensitive and should be protected from light.
Storage:	RT,4 °C,-20 °C
Expiry Date:	12 months

Images



Western Blotting

Image 1. Western Blot of Anti-Rat IgG (H&L) (SHEEP) Antibody . Lane M: 3 µl Molecular Ladder. Lane 1: Rat IgG whole molecule . Lane 2: Rat IgG F(c) Fragment . Lane 3: Rat IgG Fab Fragment . Lane 4: Rat IgM Whole Molecule . Lane 5: Rat Serum . All samples were reduced. Load: 50 ng per Iane. Block: ABIN925618 for 30 min at RT. Primary Antibody: Anti-Rat IgG (H&L) (SHEEP) Antibody 1:1,000 for 60 min at RT. Secondary Antibody: Anti-Sheep IgG (DONKEY) Peroxidase Conjugated Antibody 1:40,000 in ABIN925618 for 30 min at RT. Predicted/Obsevered Size: 25 and 55 kDa for Rat IgG and Serum, 25 kDa for F(c) and Fab, 78 and 25 kDa for IgM. Rat F(c) migrates slightly higher.

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