

Datasheet for ABIN6699037 Sheep anti-Mouse IgG Antibody (DyLight 800)

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Abstract:

Overview		
Quantity:	100 µg	
Target:	IgG	
Reactivity:	Mouse	
Host:	Sheep	
Clonality:	Polyclonal	
Conjugate:	DyLight 800	
Application:	Western Blotting (WB), FLISA, Fluorescence Microscopy (FM), Dot Blot (DB)	
Product Details		
Purpose:	Mouse IgG (H&L) Antibody DyLight™ 800 Conjugated	
Immunogen:	Mouse IgG whole molecule	
Isotype:	IgG	
Characteristics:	Sheep Anti-Mouse IgG Antibody DyLight™ 800 Conjugated, Sheep Anti Mouse IgG (H&L) Antibody DyLight™ 800 Conjugated,Anti-Mouse IgG DyLight 800 Antibody generated in sheep detects reactivity to Mouse IgG.	
Labeling Ratio:	2.7	
Target Details		
Target:	lgG	
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IgG Products

Target Details

Target Type:AntibodyBackground: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 opsonization 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.

Application Details

Application Notes:	FLISA_Dilution: >1:20,000
	IF_Microscopy_Dilution: >1:5,000
	Western_Blot_Dilution: >1:10,000
	Other: User Optimized
Comment:	Anti-Mouse IgG DyLight 800 Antibody has been tested by dot blot and is designed for
	immunofluorescence microscopy, fluorescence based plate assays (FLISA) and fluorescent
	western blotting. This product is also suitable for multiplex analysis, including multicolor
	imaging, utilizing various commercial platforms. The emission spectra for this DyLight $^{\scriptscriptstyle M}$
	conjugate match the principle output wavelengths of most common fluorescence
	instrumentation.
	Suggested Applications: WB
Restrictions:	For Research Use only
Handling	
Format:	Lyophilized
Reconstitution:	Reconstitution Volume: 100 µL
	Reconstitution Buffer: Restore with deionized water (or equivalent)
Concentration:	1.0 mg/mL
Buffer:	0.02 M Potassium Phosphate, 0.15 M Sodium Chloride, pH 7.2, 10 mg/mL Bovine Serum

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	Albumin (BSA) - Immunoglobulin and Protease free, 0.01 $\%$ (w/v) Sodium Azide
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	4 °C,-20 °C
Storage Comment:	Store conjugated secondary antibody at 4° C prior to restoration. For extended storage aliquot contents and freeze at -20° C or below. Avoid cycles of freezing and thawing. Centrifuge product if not completely clear after standing at room temperature. Conjugated Secondary Antibody is stable for several weeks at 4° C as an undiluted liquid. Dilute only prior to immediate use.
Expiry Date:	12 months
Publications	
Product cited in:	Becker, Plückthun: "DARPins bind their cytosolic targets after having been translocated through the protective antigen pore of anthrax toxin." in: Scientific reports , Vol. 13, Issue 1, pp. 8048, (2023) (PubMed).
	Yan, Zhu, Liang, Feng: "NE-activated β2-AR/β-arrestin 2/Src pathway mediates duodenal hyperpermeability induced by water-immersion restraint stress." in: American journal of physiology. Cell physiology , Vol. 324, Issue 1, pp. C133-C141, (2023) (PubMed).
	Navarro, Tapia-Galisteo, Martín-García, Tarín, Corbacho, Gómez-López, Sánchez-Tirado, Campuzano, González-Cortés, Yáñez-Sedeño, Compte, Álvarez-Vallina, Sanz: "TGF-β-induced IGFBP-3 is a key paracrine factor from activated pericytes that promotes colorectal cancer cell migration and invasion." in: Molecular oncology , Vol. 14, Issue 10, pp. 2609-2628, (2021) (PubMed).