

Datasheet for ABIN102262

Rabbit anti-Sheep IgG (Heavy & Light Chain) Antibody (HRP) - Preadsorbed



1	Image	2	Publications
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Overview	
Quantity:	1 mg
Target:	IgG
Binding Specificity:	Heavy & Light Chain
Reactivity:	Sheep
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	HRP
Application:	ELISA, Immunohistochemistry (IHC), Western Blotting (WB)
Product Details	
Purpose:	Sheep IgG (H&L) Antibody Peroxidase Conjugated Pre-Adsorbed
Immunogen:	Optional[Immunogen]: Sheep IgG whole molecule
lsotype:	IgG
Cross-Reactivity (Details):	Minimal crossreactivity against Human Serum Proteins Assay by immunoelectrophoresis resulted in a single precipitin arc against anti-Peroxidase, anti-Rabbit Serum, Sheep IgG and Sheep Serum. No reaction was observed against Human Serum Proteins.
Characteristics:	Anti-Sheep IgG F(ab')2 Antibody generated in rabbit recognizes the dimeric Fab portion of the sheep IgG molecule. Sheep IgG F(ab')2 is a proteolytic fragment of immunoglobulin G (IgG) obtained by limited digestion with the enzyme pepsin under controlled conditions of temperature, time and pH . F(ab')2 Molecules lack the Fc portion of IgG and therefore receptor

that bind sheep IgG F(c) will not bind sheep IgG F(ab')2 Molecules. Secondary Antibodies are

Product Details

Troduct Details	
	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-Sheep IgG F(ab')2 is conjugated to Peroxidase.
Purification:	Secondary Antibody was prepared from monospecific antiserum by immunoaffinity
	chromatography using Sheep IgG coupled to agarose beads followed by solid phase
	adsorption(s) to remove any unwanted reactivities.
Target Details	
Target:	IgG
Abstract:	IgG Products
Target Type:	Antibody
Background:	HRP Conjugated Secondary Antibody is designed for western blotting, ELISA and
	Immunohistochemistry. Horseradish peroxidase conjugated antibodies are ideal for antibody
	detection assays that require lot-to-lot consistency.
Application Details	
Application Notes:	Application Note: Anti-Sheep IgG Peroxidase Conjugated Antibody has been tested by ELISA
	and is suitable for immunoblotting (western or dot blot), ELISA, immunoperoxidase electron
	microscopy and immunohistochemistry as well as other peroxidase-antibody based enzymatic
	assays. Immunohistochemistry Dilution: 1:300 - 1:1,500 Western Blot Dilution: 1:500 - 1:3,000
	ELISA Dilution: 1:5,000 - 1:22,000 Other: User Optimized
Restrictions:	For Research Use only
Handling	
Format:	Lyophilized
Reconstitution:	Reconstitution Buffer: Restore with deionized water (or equivalent), Reconstitution Volume: 1.0
	mL
Concentration:	1.0 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) Gentamicin Sulfate. Do NOT add Sodium Azide!

Handling

Preservative:	Gentamicin sulfate
Precaution of Use:	This product contains Gentamicin sulfate: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	4 °C,-20 °C
Storage Comment:	Store 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. This product is stable for several weeks at 4° C as an undiluted liquid. Dilute only prior to immediate use.
Expiry Date:	12 months

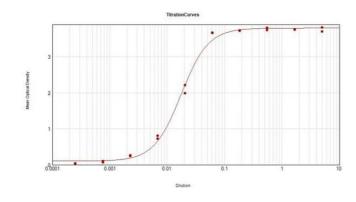
Publications

Product cited in:

Bissoon-Haqqani, Moyana, Jonker, Maroun, Birnboim: "Nuclear expression of thymidylate synthase in colorectal cancer cell lines and clinical samples." in: **The journal of histochemistry and cytochemistry : official journal of the Histochemistry Society**, Vol. 54, Issue 1, pp. 19-29, (2006) (PubMed).

Cripe, Gelfanov, Smith, Spigel, Phillips, Gabig, Jung, Fyffe, Hartman, Kneebone, Mercola, Burgess, Boswell: "Role for c-jun N-terminal kinase in treatment-refractory acute myeloid leukemia (AML): signaling to multidrug-efflux and hyperproliferation." in: **Leukemia**, Vol. 16, Issue 5, pp. 799-812, (2002) (PubMed).

Images



ELISA

Image 1. ELISA Results of Rabbit Anti-Sheep IgG Antibody Peroxidase Conjugate (Min X Human Serum Proteins) tested against purified Sheep IgG Min X Hu HRP. Each well was coated in duplicate with 1.0 μ g of Sheep IgG (p/n 013-0102). The working dilution is 1:57000. The starting dilution of antibody was 5 μ g/mL and the X-axis represents the Log10 of a 3-fold dilution. This titration is a 4-parameter curve fit where the IC50 is defined as the titer of the

antibody.