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Goat anti-Rat IgG (Heavy & Light Chain) Antibody (HRP)

2 Images

Overview

Clonality:

2

Publications



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1 mg
IgG
Heavy & Light Chain
Rat
Goat

Conjugate: HRP

Application: ELISA, Immunohistochemistry (IHC), Western Blotting (WB), Dot Blot (DB), Immunoelectron

Microscopy (IEM)

Polyclonal

Product Details

Immunogen:	Anti-Rat IgG whole molecule was produced by repeated immunization with Rat IgG whole molecule in goat.
Isotype:	IgG
Specificity:	IgG (H&L)
Characteristics:	Anti-Rat IgG whole molecule antibody generated in goat detects specifically Rat IgG whole molecule. This secondary antibody anti-Rat is ideal for investigators who routinely perform ELISA, Sandwich ELISA, titration assays, western-blot, immunoprecipitation and more generally immunoassays.

Concentration Definition: by UV absorbance at 280 nm

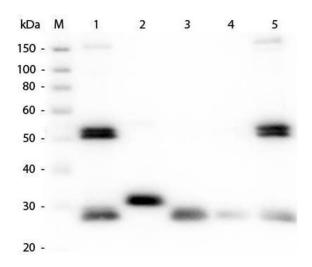
Target Details

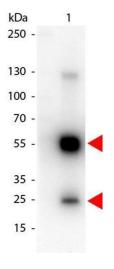
Target:	IgG
Abstract:	IgG Products
Target Type:	Antibody
Background:	Synonyms: Rat Antibody, Gt-a-Rat IgG, Rat Antibody in Goat, Rat Secondary Antibody
Application Details	
Application Notes:	Suitable for immunoblotting (western or dot blot), ELISA, immunoperoxidase electron
	microscopy and immunohistochemistry as well as other peroxidase-antibody based enzymatic assays requiring lot-to-lot consistency.
Restrictions:	For Research Use only
Handling	
Format:	Lyophilized
Reconstitution:	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
Preservative:	Gentamicin sulfate
Handling Advice:	Do NOT add Sodium Azide! Use of Sodium Azide will inhibit enzyme activity of horseradish peroxidase.
Storage:	4 °C
Publications	
Product cited in:	Mousa, Alhumaydhi, Abdellatif, Abdulmonem, AlKhowailed, Alsagaby, Al Rugaie, Alnuqaydan,
	Aljohani, Aljasir, Alwashmi, Soliman, Yosof, Elsheikh, Babiker, Alsuhaibani, Hegazy, Seleem: "
	Curcumin and ustekinumab cotherapy alleviates induced psoriasis in rats through their
	antioxidant, anti-inflammatory, and antiproliferative effects." in: Cutaneous and ocular
	toxicology , pp. 1-10, (2021) (PubMed).
	Schmitt, Funk, Blum, Asan, Andersen, Rülicke, Sendtner, Buchner: "Initial characterization of a

Syap1 knock-out mouse and distribution of Syap1 in mouse brain and cultured motoneurons."

in: Histochemistry and cell biology, Vol. 146, Issue 4, pp. 489-512, (2017) (PubMed).

Images





Western Blotting

Image 1. Western Blot of Anti-Rat IgG (H&L) (GOAT) Antibody (Min X Bv Ch Gt GP Ham Hs Hu Ms Rb & Sh Serum Proteins) . 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) (GOAT) Antibody (Min X Bv Ch Gt GP Ham Hs Hu Ms Rb & Sh Serum Proteins) 1:1,000 for 60 min at RT. Secondary Antibody: Anti-Goat 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.

Western Blotting

Image 2. Western Blot of Peroxidase conjugated Goat anti-Rat IgG antibody. Lane 1: Rat IgG. Lane 2: none. Load: 50 ng per lane. Primary antibody: none. Secondary antibody: Peroxidase rat secondary antibody at 1:1,000 for 60 min at RT. Block: ABIN925618 for 30 min at RT. Predicted/Observed size: 55 kDa, 28 kDa for Rat IgG. Other band(s): none.