

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

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Datasheet for KCB003 HRP Western Blot Anti-Rabbit IgG Antibody

Overview

Description:	HRP Western Blot Anti-Rabbit IgG Antibody - KCB003
Item No.:	KCB003
Size:	100 µg
Applications:	WB
Host Species:	Goat

Product Details

Background:	Anti-Rabbit IgG peroxidase conjugated antibody generated in goat detects specifically rabbit IgG.
Synonyms:	Anti-Rabbit IgG Peroxidase Conjugated Antibody for Western Blot
Host Species:	Goat
Conjugate:	Peroxidase (HRP)
Clonality:	Polyclonal
Detection Kit Type:	Chemiluminescent Western Blot Kit

Target Details

Purity/Specificity:HRP Western Blot Anti-Rabbit IgG Antibody was prepared from monospecific antiserum by
immunoaffinity chromatography using Rabbit IgG coupled to agarose beads followed by solid
phase adsorption(s) to remove any unwanted reactivities. Assay by immunoelectrophoresis
resulted in a single precipitin arc against anti-Peroxidase, anti-Goat Serum, Rabbit IgG and
Rabbit Serum. No reaction was observed against Human Serum Proteins.

Application Details

Tested Applications:	WB
Application Note:	HRP Western Blot Anti-Rabbit IgG Antibody is specifically designed for immunoblotting used with Chemiluminescent Western Blot Kit (p/n KCA003) with Rabbit Primary Antibody.



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Assay Dilutions:	All assays should be optimized by the user. Recommended dilutions (if any) may be
	listed below.

Formulation

Concentration:	1.0 mg/ml by UV absorbance at 280 nm
Buffer:	0.02 M Potassium Phosphate, 0.15 M Sodium Chloride, pH 7.2
Preservative:	0.01% (w/v) Gentamicin Sulfate. Do NOT add Sodium Azide!

Shipping & Handling

Shipping Condition:	Wet Ice
Storage Condition:	See kit insert for complete instructions.
Expiration:	See kit insert for complete instructions.

Images



Western Blot

(A) LoVo cells were either exposed to PTH α (30 μ M) or transfected with non-specific siRNA (ns-siRNA) or p14ARF specific siRNA. Cells were treated with 2.5 μ M oxaliplatin 8 h after siRNA or 1 h after PTH α treatment. 120 h upon oxaliplatin exposure, the expression of p14ARF, p21CIP1, and p53, as well as the phosphorylation of p53 at Ser15 was measured by immunodetection. HRP conjugated goat anti-mouse (p/n KCB002) and HRP conjugated goat anti-rabbit (p/n KCB003) were used. Fig 7. PMID: 33922007.

References

- Tomicic MT et al. Oxaliplatin-Induced Senescence in Colorectal Cancer Cells Depends on p14 ARF-Mediated Sustained p53 Activation. *Cancers (Basel)*. (2021)
- Schwarzenbach C et al. Targeting c-IAP1, c-IAP2, and Bcl-2 Eliminates Senescent Glioblastoma Cells Following Temozolomide Treatment. *Cancers (Basel)*. (2021)

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