abcam

Product datasheet

Goat Anti-Mouse IgG H&L ab182017

2 References 画像数 2

製品の概要

製品名 Goat Anti-Mouse IgG H&L

由来種Goatターゲット生物種Mouse

特異性 The antibody reacts with mouse immunoglobulins of all classes. Cross-reactions as determined

by ELISA (see figure below): Chicken lgY, less than 2%. Human lgG, less than 6%. Rabbit lgG,

less than 7%. Rat IgG, less than 47%.

アプリケーション 適用あり: ELISA, WB, IHC-P, IP

免疫原 The details of the immunogen for this antibody are not available.

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

Avoid freeze / thaw cycle.

バッファー pH: 7.40

Preservative: 0.02% Sodium azide

Constituent: PBS

精製度 Affinity purified

特記事項(精製) Immunogen affinity purified - This antibody was isolated by affinity chromatography using antigen

coupled to agarose beads.

ポリ/モノ ポリクローナル

アイソタイプ IgG

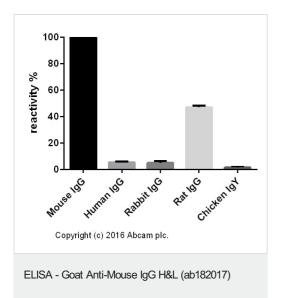
アプリケーション

The Abpromise guarantee Abpromise保証は、次のテスト済みアプリケーションにおけるab182017の使用に適用されます

アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

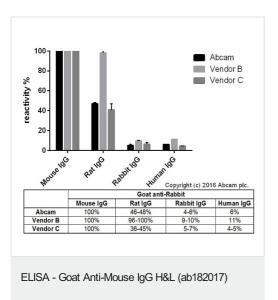
アプリケーション	Abreviews	特記事項
ELISA		1/20000 - 1/200000.
WB		Use at an assay dependent concentration.
IHC-P		Use at an assay dependent concentration.
IP		Use at an assay dependent concentration.

画像



Cross-reactivity of the polyclonal secondary antibody ab182017 was tested using a sandwich ELISA approach. The wells were coated with the indicated IgG standards at 1 μ g/ml (50 μ l/well) and incubated overnight at 4°C, followed by a 5% BSA blocking step for 2h at RT. ab182017 was then added starting at 1 μ g/ml and gradually diluted 1/4 (50 μ l/well), followed by incubation for 2h. For the detection Donkey anti-Goat IgG H&L (HRP) (ab6885) was used at 1/10,000 dilution (50 μ l/well), followed by incubation for 1h at RT.

Fot the batch tested, ab182017 showed a cross-reactivity below 2% towards Chicken IgY, 6% towards Human IgG, 7% towards Rabbit IgG and 47% towards Rat IgG.



Cross-reactivity of Goat anti-Mouse IgG H&L (ab182017) and Goat anti-Mouse IgG H&L obtained from two different vendors was tested using a sandwich ELISA approach. The wells were coated with the indicated IgG standards (Rabbit, Human, Mouse and Rat) at 1 μ g/ml (50 μ l/well) and incubated overnight at 4°C, followed by a 5% BSA blocking step for 2h at RT. Secondary antibodies were then added starting at 1 μ g/ml and gradually diluted 1/4 (50 μ l/well), followed by incubation for 2h. For the detection Donkey anti-Goat IgG H&L (HRP) (ab6885) was used at 1/10,000 dilution (50 μ l/well), followed by incubation for 1h at RT. This data is from a representative dilution.

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