abcam

Product datasheet

Goat Anti-Rabbit IgG H&L (Alexa Fluor® 790) ab175781

22 References 3 图像

概述

产品名称 山羊抗兔lgG H&L (Alexa Fluor® 790)

宿主 Goat **靶标种属** Rabbit

经测试应用 适用于: WB

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

偶联物 Alexa Fluor® 790. Ex: 782nm, Em: 805nm

性能

形式 Liquid

存放说明 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

Avoid freeze / thaw cycle. Stable for 12 months at -20°C. Store In the Dark.

存储溶液 Preservative: 0.02% Sodium azide

Constituents: 23% Glycerol (glycerin, glycerine), PBS, 1% BSA

纯**度** Immunogen affinity purified

克隆 多克隆

常规说明

We batch test Goat Anti-Rabbit IgG H&L (Alexa Fluor® 790), ab175781 in fluorescent WB.

Although we don't batch test for ICC, ELISA, IHC-Fr or Flow cytometry customers have had

success using Goat Anti-Rabbit IgG H&L (Alexa Fluor® 790), ab175781 in these applications.

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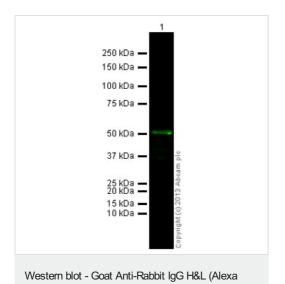
应用

The Abpromise guarantee Abpromise™承诺保证使用ab175781于以下的经测试应用

"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
WB		1/10000.

图片



Fluor® 790) (ab175781)

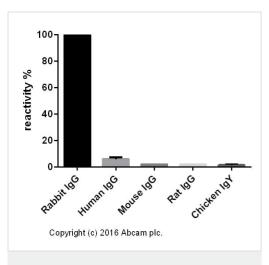
Anti-alpha Tubulin antibody - Microtubule Marker ($\underline{ab18251}$) at 1 $\mu g/ml$ + HeLa (Human epithelial carcinoma cell line) Whole Cell Lysate ($\underline{ab27252}$) at 10 μg

Secondary

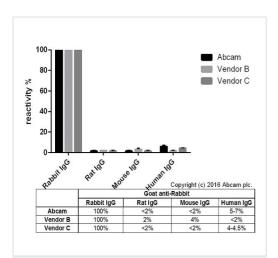
Goat Anti-Rabbit IgG H&L (Alexa Fluor® 790) (ab175781) at 1/10000 dilution

Observed band size: 50 kDa

This blot was produced using a 4-12% Bis-tris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 5% Milk before being incubated with **ab18251** overnight at 4°C. Antibody binding was detected using ab175781 at a 1:10,000 dilution for 1hr at room temperature and then imaged using the Licor Odyssey CLx.



ELISA - Goat Anti-Rabbit IgG H&L (Alexa Fluor® 790) (ab175781)



ELISA - Goat Anti-Rabbit IgG H&L (Alexa Fluor® 790) (ab175781)

Cross-reactivity of the polyclonal secondary antibody <u>ab182016</u> 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. <u>ab182016</u> 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) (<u>ab6885</u>) was used at 1/10,000 dilution (50 μ l/well), followed by incubation for 1h at RT.

For the batch tested, <u>ab182016</u> showed a cross-reactivity of 5-7% towards Human IgG and below 2% towards Mouse IgG, Rat IgG and Chicken IgY.

This data was developed using the unconjugated antibody (ab182016).

Cross-reactivity of Goat anti-Rabbit IgG H&L (ab182016) and Goat anti-Rabbit 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 µg/ml (50 µl/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 µg/ml and gradually diluted 1/4 (50 µl/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 µl/well), followed by incubation for 1h at RT. This data is from a representative dilution.

This data was developed using the unconjugated antibody (ab182016).

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