

# Goat Anti-Rabbit IgG H&L (Alexa Fluor® 680) preadsorbed ab186696

5 References [画像数 1](#)

### 製品の概要

製品名	Goat Anti-Rabbit IgG H&L (Alexa Fluor® 680) preadsorbed
由来種	Goat
ターゲット生物種	Rabbit
特異性	By immunoelectrophoresis and ELISA this antibody reacts specifically with rabbit IgG and with light chains common to other rabbit immunoglobulins. No antibody was detected against non-immunoglobulin serum proteins. Reduced cross-reactivity to bovine, chicken, horse, human, mouse, pig, and rat IgG was detected. This antibody may cross react with IgG from other species.
アプリケーション	適用あり: WB
吸着処理血清	Chicken, Cow, Horse, Human, Mouse, Pig, Rat <a href="#">more details</a>
免疫原	The details of the immunogen for this antibody are not available.
標識	Alexa Fluor® 680. Ex: 679nm, Em: 702nm

### 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. Stable for 12 months at -20°C. Store In the Dark.
バッファー	Preservative: 0.02% Sodium azide Constituents: 23% Glycerol (glycerin, glycerine), PBS, 1% BSA
精製度	Immunogen affinity purified
特記事項 (精製)	Antiserum was cross adsorbed using bovine, chicken, horse, human, mouse, pig and rat immunosorbents to remove cross reactive Antibodies. The antibody to rabbit IgG was isolated by affinity chromatography using antigen coupled to agarose beads.
ポリ/モノ	ポリクローナル
特記事項	We batch test Goat Anti-Rabbit IgG H&L (Alexa Fluor® 680) preadsorbed, ab186696 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® 680) preadsorbed, ab186696 in

these applications.

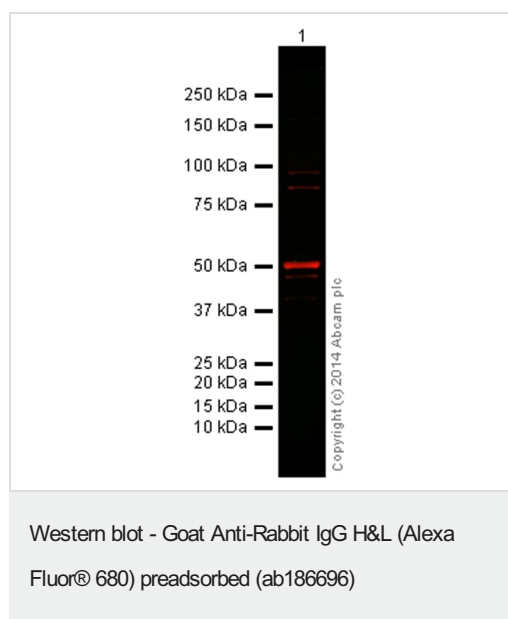
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## アプリケーション

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アプリケーション	Abreviews	特記事項
WB		1/10000.

## 画像



Anti-beta Tubulin antibody - Loading Control (**ab6046**) at 1 µg/ml +  
HeLa (Human epithelial carcinoma cell line) Whole Cell Lysate at  
10 µg

### Secondary

Goat Anti-Rabbit IgG H&L (Alexa Fluor® 680) preadsorbed  
(ab186696) 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 **ab6046** overnight at 4°C. Antibody binding was detected using ab186696 at a 1:10,000 dilution for 1hr at room

temperature and then imaged using the Licor Odyssey CLx.

**Please note:** All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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