


Anti-Golgin 97 antibody ab84340

★★★★★ [6 Abreviews](#) [18 References](#) [画像数 6](#)

製品の概要

製品名	Anti-Golgin 97 antibody
製品の詳細	Rabbit polyclonal to Golgin 97
由来種	Rabbit
アプリケーション	適用あり: ICC/IF, IHC-P
種交差性	交差種: Human 交差が予測される動物種: Mouse 
免疫原	Synthetic peptide corresponding to Human Golgin 97 aa 750 to the C-terminus (C terminal) conjugated to keyhole limpet haemocyanin. (Peptide available as ab34458)
ポジティブ・コントロール	ICC/IF: HeLa, HepG2, MCF7 and hCMEC/D3 cells. IHC-P: Human breast carcinoma tissue. IHC-P: Normal Human Pancreas tissue.
特記事項	<p>ab84340 is raised using the same immunogen as ab33701. Batches sold under ab84340 have passed QC testing in ICC/IF and IHC. In our hands, this product did not work in WB. For further information please contact our Customer Services team.</p> <p>The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets your needs before purchasing.</p> <p>If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be found below, along with publications, customer reviews and Q&As</p>

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 or -80°C. Avoid freeze / thaw cycle.
バッファー	pH: 7.40 Preservative: 0.02% Sodium azide Constituent: PBS

Batches of this product that have a concentration < 1mg/ml may have BSA added as a stabilising

agent. If you would like information about the formulation of a specific lot, please contact our scientific support team who will be happy to help.

精製度	Immunogen affinity purified
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

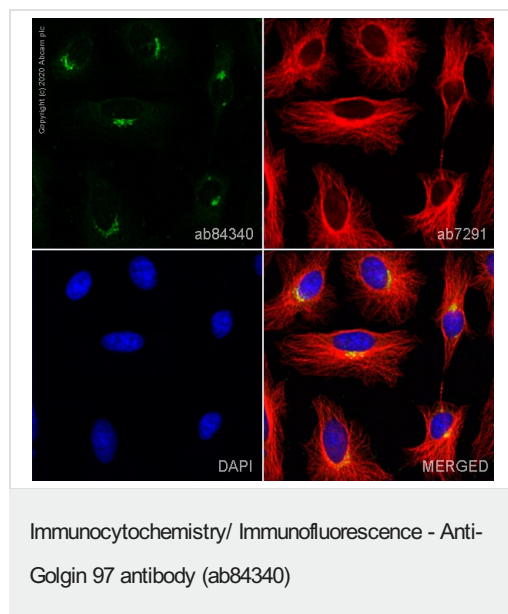
The Abpromise guarantee **Abpromise保証は、次のテスト済みアプリケーションにおけるab84340の使用に適用されます**
アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
ICC/IF	★★★★★ (4)	Use a concentration of 5 µg/ml. Suitable for cells that have been fixed with 100% Methanol (5 mins) or 4% PFA (10 mins).
IHC-P	★★★★★ (1)	Use a concentration of 5 µg/ml. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.

ターゲット情報

機能	Probably involved in maintaining Golgi structure.
配列類似性	Contains 1 GRIP domain.
ドメイン	Extended rod-like protein with coiled-coil domains.
細胞内局在	Golgi apparatus > Golgi stack membrane.

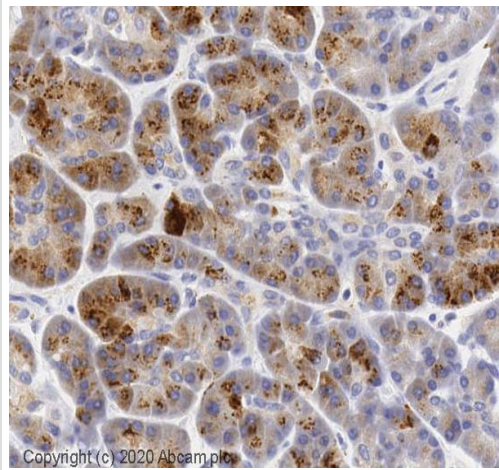
画像



ab84340 staining Golgin 97 in HeLa cells. The cells were fixed with 100% Methanol (5min), permeabilized with 0.1%PBS-Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1%PBS-Tween for 1h. The cells were then incubated overnight at +4°C with ab84340 at 5µg/ml and **ab7291**, Mouse monoclonal [DM1A] to alpha Tubulin - Loading Control, at 1/1000 dilution. Cells were then incubated with **ab150081**, Goat polyclonal Secondary Antibody to Rabbit IgG - H&L (Alexa Fluor® 488), pre-adsorbed (shown in green) and **ab150120**, Goat polyclonal Secondary Antibody to Mouse IgG - H&L (Alexa Fluor® 594), pre-adsorbed at 1/1000 dilution (shown in pseudocolor red). Nuclear DNA was labelled with DAPI (shown in blue).

Image was taken with a confocal microscope (Leica-Microsystems,

TCS SP8).



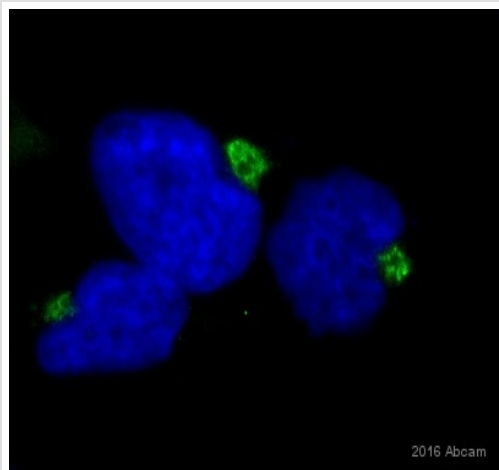
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Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Golgin 97 antibody (ab84340)

IHC image of Golgin 97 antibody staining in a section of formalin-fixed paraffin-embedded normal human pancreas* performed on a Leica BOND™ system using the standard protocol. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20mins. The section was then incubated with ab84340, 1ug/ml, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.

**Tissue obtained from the Human Research Tissue Bank, supported by the NIHR Cambridge Biomedical Research Centre*

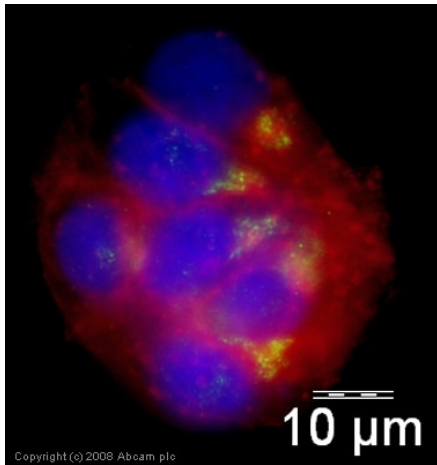


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Immunocytochemistry/ Immunofluorescence - Anti-Golgin 97 antibody (ab84340)

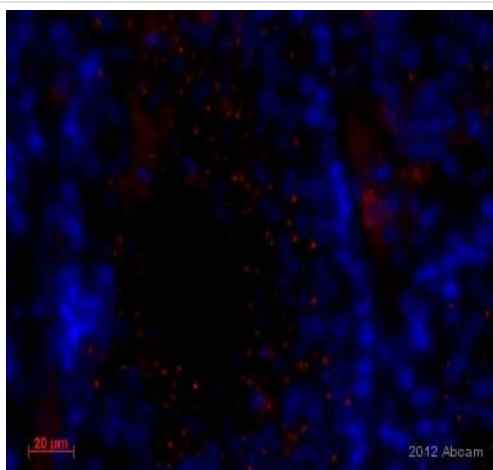
This image is courtesy of an anonymous Abreview.

hCMEC/D3 cells stained for Golgin 97 (green) using ab84340 at 1/100 dilution in ICC/IF, followed by Alexa Fluor 488® conjugated Goat Anti-Rabbit IgG.



Immunocytochemistry/ Immunofluorescence - Anti-Golgin 97 antibody (ab84340)

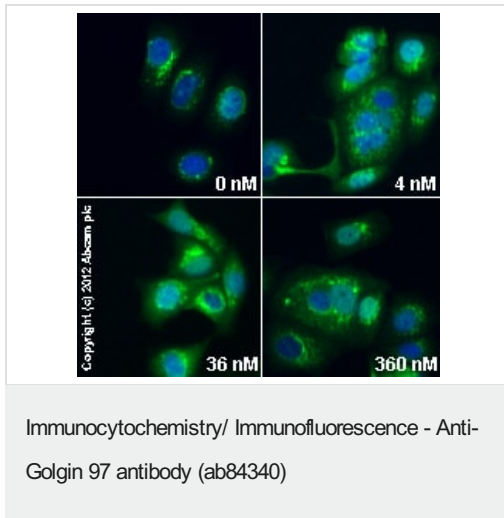
ICC/IF image of ab84340 stained human HepG2 cells. The cells were methanol fixed (5 min), permeabilised in 0.1% PBS-Tween (20 min) and incubated with the antibody (ab84340, 5μg/ml) for 1h at room temperature. 1%BSA / 10% normal goat serum / 0.3M glycine was used to block non-specific protein-protein interactions. The secondary antibody (green) was Alexa Fluor® 488 goat anti-rabbit IgG (H+L) used at a 1/1000 dilution for 1h. Alexa Fluor® 594 WGA was used to label plasma membranes (red). DAPI was used to stain the cell nuclei (blue).



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Golgin 97 antibody (ab84340)

ab84340 staining Golgin 97 in murine testis tissue by Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections). Tissue was fixed with formaldehyde and a heat mediated antigen retrieval step was performed using 0.01M sodium citrate. Samples were then blocked using 10%FCS, 0.5% BSA, 1% Triton X-100, 88.5% PBS X1 and then incubated with ab84340 at a 1/100 dilution for 10 hours at 4°C. The secondary used was an Alexa-Fluor 594 conjugated goat anti-rabbit IgG used at a 1/200 dilution.

Image courtesy of Yoav Elkis by Abreview.



ab84340 staining Golgin-97 in MCF7 cells treated with brefeldin A (**ab120299**), by ICC/IF. Increase in Golgin-97 expression correlates with increased concentration of brefeldin A, as described in literature.

The cells were incubated at 37°C for 1.5 h in media containing different concentrations of **ab120299** (brefeldin A) in DMSO, fixed with 4% formaldehyde for 10 minutes at room temperature and blocked with PBS containing 10% goat serum, 0.3 M glycine, 1% BSA and 0.1% tween for 2h at room temperature. Staining of the treated cells with ab84340 (5 µg/ml) was performed overnight at 4°C in PBS containing 1% BSA and 0.1% tween. A DyLight 488 goat anti-rabbit polyclonal antibody (**ab96899**) at 1/250 dilution was used as the secondary antibody. Nuclei were counterstained with DAPI and are shown in blue.

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