abcam

Product datasheet

Anti-SKP2 antibody [EPR3305(2)] ab183039



★★★★★ 1 Abreviews 10 References

画像数 6

製品の概要

製品名 Anti-SKP2 antibody [EPR3305(2)]

製品の詳細 Rabbit monoclonal [EPR3305(2)] to SKP2

由来種 Rabbit

アプリケーション 適用あり: Flow Cyt (Intra), ICC/IF, IHC-P, WB

種交差性 交差種: Mouse, Rat, Human

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

ポジティブ・コントロール WB: C6, NIH/3T3, F9, A549, MDA-MB-231, Jurkat, MCF7, HepG2, and NCCIT cell lysates. ICC:

293T cells

特記事項 This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity

- Long-term security of supply

- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

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 long

term. Avoid freeze / thaw cycle.

バッファー pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA

精製度 Protein A purified

ポリモノ モノクローナル

クローン名 EPR3305(2)

ΙgG アイソタイプ

The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab183039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
Flow Cyt (Intra)		1/70. ab172730 - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody.
ICC/IF	★★★ ☆☆ <u>(1)</u>	1/100.
IHC-P		1/50. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
WB		1/200. Detects a band of approximately 48 kDa (predicted molecular weight: 48 kDa).

ターゲット情報

機能

Substrate recognition component of a SCF (SKP1-CUL1-F-box protein) E3 ubiquitin-protein ligase complex which mediates the ubiquitination and subsequent proteasomal degradation of target proteins involved in cell cycle progression, signal transduction and transcription. Specifically recognizes phosphorylated CDKN1B/p27kip and is involved in regulation of G1/S transition. Degradation of CDKN1B/p27kip also requires CKS1. Recognizes target proteins ORC1, CDT1, RBL2, MLL, CDK9, RAG2, FOXO1A, UBP43, and probably MYC, TOB1 and TAL1. Degradation of TAL1 also requires STUB1. Recognizes CDKN1A in association with CCNE1 or CCNE2 and CDK2.

パスウェイ

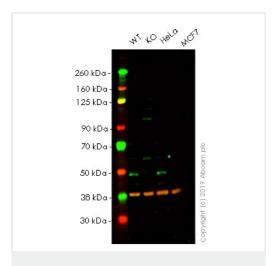
Protein modification; protein ubiquitination.

配列類似性

Contains 1 F-box domain.

Contains 9 LRR (leucine-rich) repeats.

画像



Western blot - Anti-SKP2 antibody [EPR3305(2)] (ab183039)

All lanes : Anti-SKP2 antibody [EPR3305(2)] (ab183039) at 1/200 dilution

Lane 1: Wild-type HAP1 whole cell lysate

Lane 2: SKP2 knockout HAP1 whole cell lysate

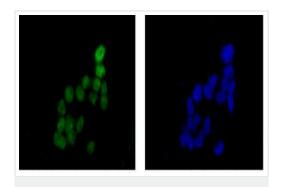
Lane 3 : HeLa whole cell lysate
Lane 4 : MCF7 whole cell lysate

Lysates/proteins at 20 µg per lane.

Predicted band size: 48 kDa

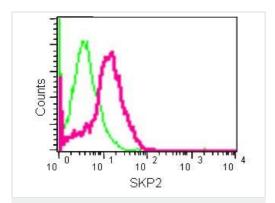
Lanes 1 - 4: Merged signal (red and green). Green - ab183039 observed at 52 kDa. Red - loading control, **ab8245**, observed at 37 kDa.

ab183039 was shown to recognize in wild-type HAP1 cells as signal was lost at the expected MW in SKP2 knockout cells. Additional cross-reactive bands were observed in the wild-type and knockout cells. Wild-type and SKP2 knockout samples were subjected to SDS-PAGE. The membrane was blocked with 3% NF Milk. Ab183039 and ab8245 (Mouse anti-GAPDH loading control) were incubated overnight at 4°C at 1/200 dilution and 1/20000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preabsorbed ab216773 and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed ab216776 secondary antibodies at 1/20000 dilution for 1 hour at room temperature before imaging.



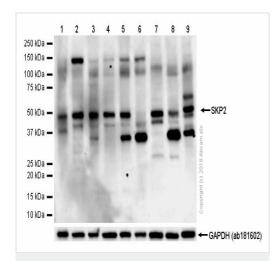
Immunocytochemistry/ Immunofluorescence - Anti-SKP2 antibody [EPR3305(2)] (ab183039)

Immunofluorescenct analysis of 4% paraformaldehyde fixed 293T cells labeling SKP2 with ab183039 at 1/100 followed by Goat anti rabbit lgG(Alexa Fluor® 488) at 1/200 (green). Cells were counter stained with Dapi (blue).



Flow Cytometry (Intracellular) - Anti-SKP2 antibody [EPR3305(2)] (ab183039)

Intracellular flow cytometric analysis of 2% paraformaldehyde fixedHeLa cells labeling SKP2 with ab183039 at 1/70 followed byGoat anti rabbit lgG (FITC) at 1/150.Rabbit monoclonal lgG was used aslsotype control.



Western blot - Anti-SKP2 antibody [EPR3305(2)] (ab183039)

All lanes : Anti-SKP2 antibody [EPR3305(2)] (ab183039) at 1/200 dilution

Lane 1 : NCCIT (Human pluripotent embryonic carcinoma epithelial cell) whole cell lysates with 5% NFDM/TBST

Lane 2: HepG2 (Human hepatocellular carcinoma epithelial cell) whole cell lysates with 5% NFDM/TBST

Lane 3 : MCF7 (Human breast adenocarcinoma epithelial cell) whole cell lysates with 5% NFDM/TBST

Lane 4: Jurkat (Human T cell leukemia T lymphocyte) whole cell lysates with 5% NFDM/TBST

Lane 5 : MDA-MB-231 (Human breast adenocarcinoma epithelial cell) whole cell lysates with 5% NFDM/TBST

Lane 6: A549 (Human lung carcinoma epithelial cell) whole cell lysates with 5% NFDM/TBST

Lane 7: F9 (Mouse embryonal carcinoma epithelial cell) whole cell lysates with 5% NFDM/TBST

Lane 8 : NIH/3T3 (Mouse embryonic fibroblast) whole cell lysates

with 5% NFDM/TBST

Lane 9 : C6 (Rat glial tumor glial cell) whole cell lysate with 5% NFDM/TBST

Lysates/proteins at 20 µg per lane.

Secondary

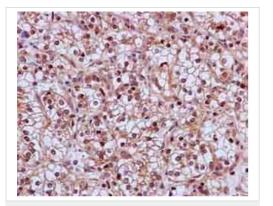
All lanes : Goat Anti-Rabbit lgG H&L (HRP) ($\underline{ab97051}$) at 1/20000

dilution

Predicted band size: 48 kDa

Exposure time: 180 seconds

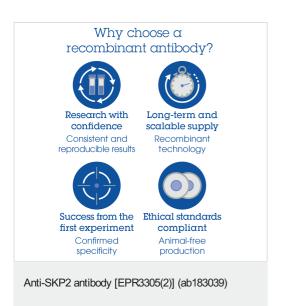
We recommend to increase the amount of samples or decrease antibody dilution to get clear bands.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-SKP2 antibody
[EPR3305(2)] (ab183039)

immunohistochemical analysis of paraffin embedded Human kidney clear cell carcinoma tissue labeling SKP2 with ab183039 at 1/50 followed by secondary staining with Ready to use HRP Polymer for Rabbit IgG and counterstained with Hematoxylin.

Perform heat mediated antigen retrieval with EDTA buffer pH 9 before commencing with IHC staining protocol.



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