abcam

Product datasheet

Anti-Calpastatin antibody [SP82] ab101684

יעלעבע RabMAb

画像数6

製品の概要

製品名 Anti-Calpastatin antibody [SP82]

製品の詳細 Rabbit monoclonal [SP82] to Calpastatin

由来種 Rabbit

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

種交差性 交差種: Human

免疫原 Synthetic peptide within Human Calpastatin aa 550-650 (C terminal). The exact sequence is

proprietary.

Database link: P20810

ポジティブ・コントロール IHC-P: Human breast carcinoma tissue; Flow Cyt (Intra): HaCaT and HeLa cells; ICC/IF: HaCaT

特記事項 This product is FOR RESEARCH USE ONLY. For commercial use, please contact

partnerships@abcam.com.

製品の特性

製品の状態 Liquid

Shipped at 4°C. Store at +4°C. Do Not Freeze. 保存方法

バッファー pH: 7.60

> Preservative: 0.1% Sodium azide Constituents: PBS, 1% BSA

精製度 Protein A/G purified

特記事項(精製) Purified from TCS by protein A/G.

ポリ/モノ モノクローナル

クローン名 SP82 アイソタイプ ΙgG

アプリケーション

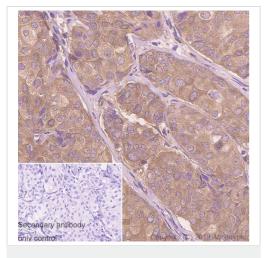
The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab101684の使用に適用されます アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
ICC/IF		Use at an assay dependent concentration.
IHC-P		1/100. Incubate for 30 minutes at RT. Antigen retrieval is recommended: boil tissue section in 10mM citrate buffer, pH 6.0 for 10 min followed by cooling at RT for 20 min.
Flow Cyt (Intra)		Use at an assay dependent concentration. <u>ab172730</u> - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody.

ターゲット情報

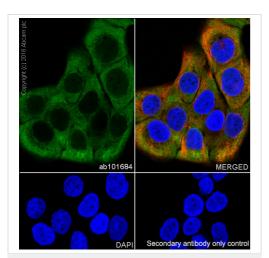
機能	Specific inhibition of calpain (calcium-dependent cysteine protease). Plays a key role in postmortem tenderization of meat and have been proposed to be involved in muscle protein degradation in living tissue.
配列類似性	Belongs to the protease inhibitor l27 (calpastatin) family.
ドメイン	Has four inhibitory domains.
翻訳後修飾	The N-terminus is blocked.

画像



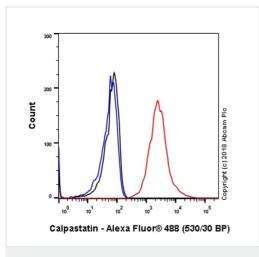
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Calpastatin antibody
[SP82] (ab101684)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of Human breast carcinoma tissue sections labeling Calpastatin with Purified ab101684 at 1/100 dilution (0.59 µg/ml). Heat mediated antigen retrieval was performed Heat mediated antigen retrieval with Citrate buffer (pH 6.0, epitope retrieval solution 1) for 20mins. Rabbit specific IHC polymer detection kit HRP/DAB (ab209101) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain.



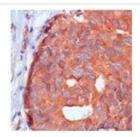
Immunocytochemistry/ Immunofluorescence - Anti-Calpastatin antibody [SP82] (ab101684)

Immunocytochemistry/ Immunofluorescence analysis of HaCaT (human skin keratinocyte) cells labeling Calpastatin with purified ab101684. Cells were fixed in 4% paraformaldehyde and permeabilized with 0.1% Triton X-100. Cells were counterstained with ab195889 Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor[®] 594). Goat anti rabbit IgG (Alexa Fluor[®] 488, ab150077) was used as the secondary antibody. DAPI was used as nuclear counterstain. PBS instead of the primary antibody was used as the secondary antibody only control.



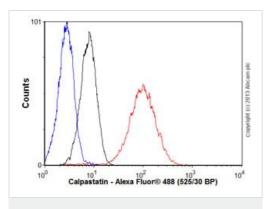
Flow Cytometry (Intracellular) - Anti-Calpastatin antibody [SP82] (ab101684)

Flow cytometry analysis of HaCaT (human skin keratinocyte) labeling Calpastatin with purified ab101684 (red). Cells were fixed with 4% paraformaldehyde and permeabilised with 90% methanol. Goat anti rabbit lgG (Alexa Fluor® 488, <u>ab150077</u>) was used as a secondary antibody. Isotype control - Rabbit monoclonal lgG (<u>ab172730</u>) (black). Unlableled control - Unlabelled cells (blue).



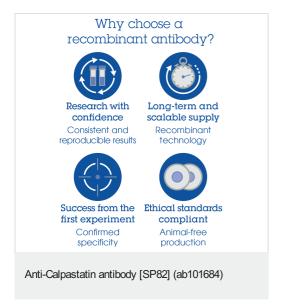
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Calpastatin antibody
[SP82] (ab101684)

Immunohistochemical analysis of Calpastatin in formalin-fixed, paraffin-embedded Human breast carcinoma using ab101684 at 1/200 dilution.



Flow Cytometry (Intracellular) - Anti-Calpastatin antibody [SP82] (ab101684)

Overlay histogram showing HeLa cells stained with ab101684 (red line). The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab101684) for 30 min at 22°C. The secondary antibody used was Alexa Fluor® 488 goat anti-rabbit IgG (H&L) (ab150077) for 30 min at 22°C. Isotype control antibody (black line) was rabbit IgG (monoclonal) (1µg/1x10⁶ cells) used under the same conditions. Unlabelled sample (blue line) was also used as a control. Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 525/30 bandpass filter.



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