


Anti-Proteasome 26S S3/PSMD3 antibody ab3316

1 References **画像数 4**

製品の概要

製品名	Anti-Proteasome 26S S3/PSMD3 antibody
製品の詳細	Rabbit polyclonal to Proteasome 26S S3/PSMD3
由来種	Rabbit
特異性	Detects proteasome 26S subunit S3.
アプリケーション	適用あり: Flow Cyt, ICC/IF
種交差性	交差種: Mouse, Human 交差が予測される動物種: Cow, Drosophila melanogaster, Non human primates 
免疫原	Synthetic peptide corresponding to Human Proteasome 26S S3/PSMD3 aa 513-534. Sequence: EREQQDLEFAKEMAEDDDDSFP

 [Run BLAST with](#)

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特記事項

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.

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

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	Constituents: 0.1% BSA, 99% PBS
精製度	Immunogen affinity purified
ポリ/モノ	ポリクローナル
アイソタイプ	IgG

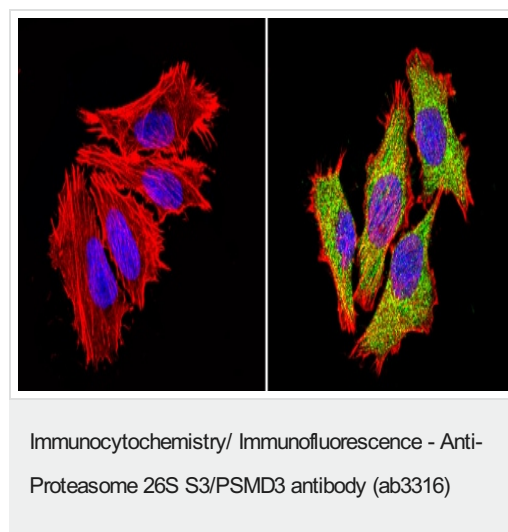
The Abpromise guarantee Abpromise保証は、 次のテスト済みアプリケーションにおけるab3316の使用に適用されます
アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
Flow Cyt		Use a concentration of 1 - 20 µg/ml.
ICC/IF		1/50 - 1/500.

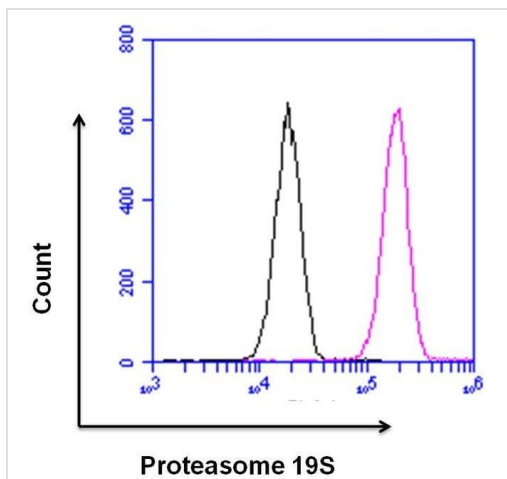
ターゲット情報

機能	Acts as a regulatory subunit of the 26 proteasome which is involved in the ATP-dependent degradation of ubiquitinated proteins.
配列類似性	Belongs to the proteasome subunit S3 family. Contains 1 PCI domain.

画像

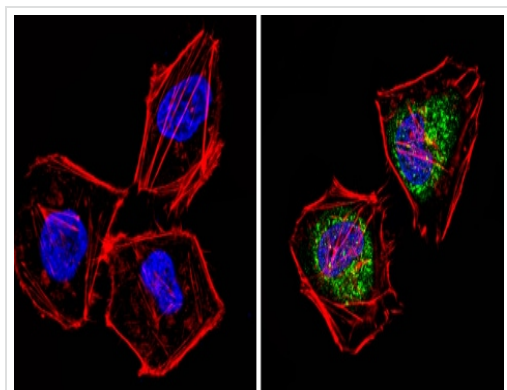


Immunocytochemistry/Immunofluorescence analysis of Proteasome 26S S3/PSMD3 (green) showing staining in the cytoplasm and nucleus of A549 cells (right) compared to a negative control without primary antibody (left). Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were incubated with ab3316 in 3% BSA-PBS at a dilution of 1:100 and incubated overnight at 4°C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight-conjugated secondary antibody in PBS at room temperature in the dark. F-actin (red) was stained with a fluorescent red phalloidin and nuclei (blue) were stained with Hoechst or DAPI. Images were taken at a magnification of 60x.



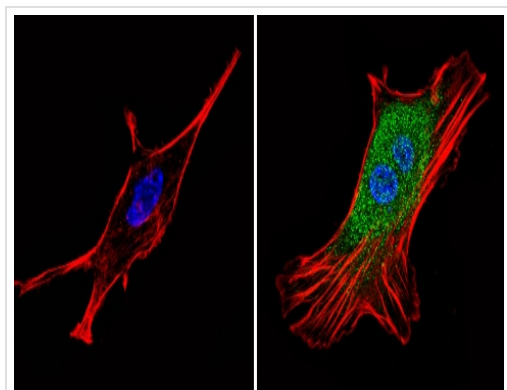
Flow Cytometry - Anti-Proteasome 26S S3/PSMD3 antibody (ab3316)

Flow Cytometry analysis of HeLa cells labeling Proteasome 26S S3/PSMD3 with ab3316 (Pink) or a rabbit IgG isotype control (Black) 10 µg/mL. Goat anti-Rabbit IgG (H+L) Superclonal™ Alexa Fluor® 647 conjugate at a dilution of 1/50 was used as the Secondary Antibody.



Immunocytochemistry/ Immunofluorescence - Anti-Proteasome 26S S3/PSMD3 antibody (ab3316)

Immunocytochemistry/Immunofluorescence analysis of Proteasome 26S S3/PSMD3 (green) showing staining in the cytoplasm and nucleus of HeLa cells (right) compared to a negative control without primary antibody (left). Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were incubated with ab3316 in 3% BSA-PBS at a dilution of 1:100 and incubated overnight at 4°C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight-conjugated secondary antibody in PBS at room temperature in the dark. F-actin (red) was stained with a fluorescent red phalloidin and nuclei (blue) were stained with Hoechst or DAPI. Images were taken at a magnification of 60x.



Immunocytochemistry/ Immunofluorescence - Anti-Proteasome 26S S3/PSMD3 antibody (ab3316)

Immunocytochemistry/Immunofluorescence analysis of Proteasome 26S S3/PSMD3 (green) showing staining in the cytoplasm and nucleus of NIH-3T3 cells (right) compared to a negative control without primary antibody (left). Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were incubated with ab3316 in 3% BSA-PBS at a dilution of 1:100 and incubated overnight at 4°C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight-conjugated secondary antibody in PBS at room temperature in the dark. F-actin (red) was stained with a fluorescent red phalloidin and nuclei (blue) were stained with Hoechst or DAPI. Images were taken at a magnification of 60x.

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