abcam

Product datasheet

Anti-Proteasome 20S beta 6 antibody ab3331

1 References 7 图像

概述

产品名称	Anti-Proteasome 20S beta 6抗体
描述	兔多克隆抗体to Proteasome 20S beta 6
宿主	Rabbit
特异性	Detects proteasome 20S beta 6 from purified bovine and human 26S proteasome samples.
经测试应用	**适用于:** ICC/IF, WB
种属反应性	**与反应:** Mouse, Rat, Cow, Human **预测可用于:** Xenopus laevis, Zebrafish
免疫原	Synthetic peptide corresponding to Human Proteasome 20S beta 6 aa 41-58. Sequence: CRSGSAADTQAV/IADAVTY Database link: **P28072** (Peptide available as **ab4947**) Run BLAST with Run BLAST with
阳性对照	WB: HeLa, BAEC whole cell lysates, HeLa, 3T3-L1, PC-3, MCF7, A549, PANC-1, Mouse Kidney, Mouse Liver and Rat Liver nuclear enriched cell lysate. ICC/IF: A431, HeLa, NIH 3T3, BAEC cells.
常规说明	Abcam has not and does not intend to apply for the REACH Authorisation of customers' uses of products that contain European Authorisation list (Annex XIV) substances. It is the responsibility of our customers to check the necessity of application of REACH Authorisation, and any other relevant authorisations, for their intended uses. 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

性能

形式	Liquid
存放说明	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.
存储溶液	Constituents: 3% Sodium deoxycholate, 3% Triton-X-100, 0.3% Tris HCl, 15% Sodium chloride
纯度	Immunogen affinity purified
克隆	**多克隆**
同种型	IgG

应用

The Abpromise guarantee **Abpromise™承诺保证使用ab3331于以下的经测试应用**

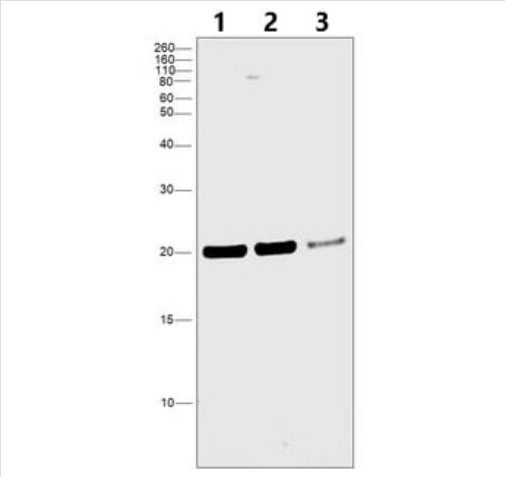
"应用说明"部分 下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
ICC/IF		1/10 - 1/100.
WB		Use a concentration of 1 µg/ml. Detects a band of approximately 25 kDa (predicted molecular weight: 25 kDa).

靶标

功能	The proteasome is a multicatalytic proteinase complex which is characterized by its ability to cleave peptides with Arg, Phe, Tyr, Leu, and Glu adjacent to the leaving group at neutral or slightly basic pH. The proteasome has an ATP-dependent proteolytic activity. This unit is responsible of the peptidyl glutamyl-like activity. May catalyze basal processing of intracellular antigens.
序列相似性	Belongs to the peptidase T1B family.
细胞定位	Cytoplasm. Nucleus.

图片



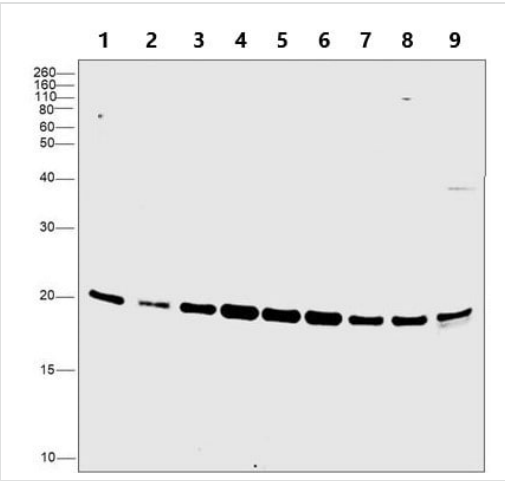
Western blot - Anti-Proteasome 20S beta 6 antibody (ab3331)

All lanes : Anti-Proteasome 20S beta 6 antibody (ab3331) at 1 µg/ml

Lane 1 : Nuclear enriched extracts from untransfected PC-3 cells
Lane 2 : Nuclear enriched extracts from non-targeting scrambled siRNA transfected PC-3 cells
Lane 3 : Nuclear enriched extracts from PSMB6 knockdown PC-3 cells

Secondary
All lanes : Goat anti-Rabbit IgG (H+L) Superclonal™ Recombinant Secondary Antibody, HRP at 1/6000 dilution

Predicted band size: 25 kDa



Western blot - Anti-Proteasome 20S beta 6 antibody (ab3331)

All lanes : Anti-Proteasome 20S beta 6 antibody (ab3331) at 1 µg/ml

Lane 1 : HeLa nuclear enriched extract lysate
Lane 2 : 3T3-L1 nuclear enriched extract lysate
Lane 3 : PC-3 nuclear enriched extract lysate
Lane 4 : MCF7 nuclear enriched extract lysate
Lane 5 : A549 nuclear enriched extract lysate
Lane 6 : PANC-1 nuclear enriched extract lysate
Lane 7 : Mouse Kidney nuclear enriched extract lysate
Lane 8 : Mouse Liver nuclear enriched extract lysate
Lane 9 : Rat Liver nuclear enriched extract lysate

Lysates/proteins at 30 µg per lane.

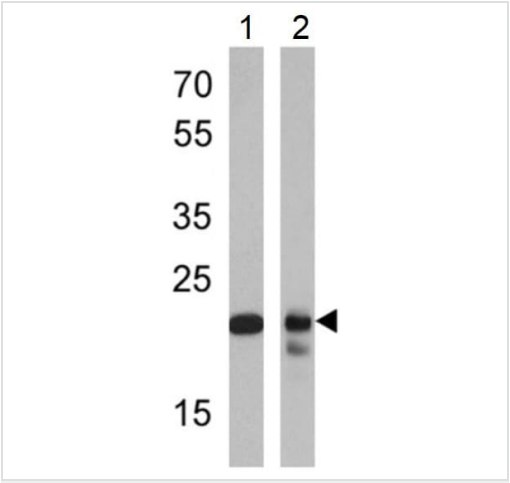
Secondary
All lanes : Goat anti-Rabbit IgG (H+L) Superclonal™ Recombinant Secondary Antibody, HRP at 1/6000 dilution

Developed using the ECL technique.

Predicted band size: 25 kDa
Observed band size: 21 kDa

Samples were electrophoresed using NuPAGE™ 12% Bis-Tris Protein Gel and the resolved proteins were then transferred onto a

Nitrocellulose membrane by iBlot® 2 Dry Blotting System. The blot was probed with the primary antibody (ab3331) and detected by chemiluminescence with secondary antibody using the iBright FL 1000. Chemiluminescent detection was performed using Novex® ECL Chemiluminescent Substrate Reagent Kit.



Western blot - Anti-Proteasome 20S beta 6 antibody (ab3331)

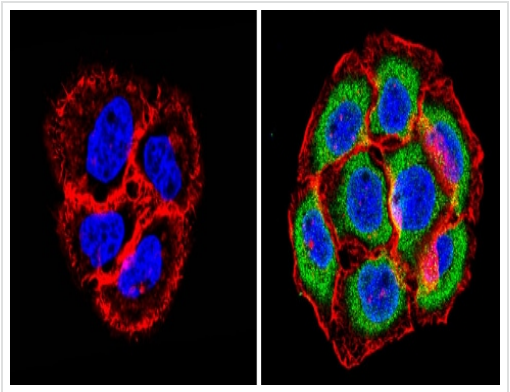
All lanes : Anti-Proteasome 20S beta 6 antibody (ab3331) at 1/1000 dilution

Lane 1 : HeLa (Human epithelial adenocarcinoma cell line) whole cell lysate
Lane 2 : BAEC (Bovine aortic endothelial cell line) whole cell lysate

Lysates/proteins at 25 µg per lane.

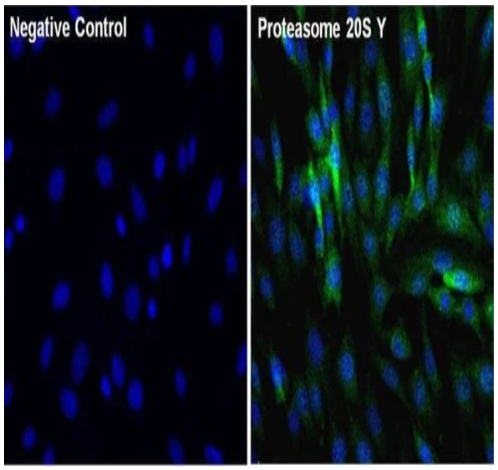
Secondary
All lanes : HRP-conjugated secondary antibody

Predicted band size: 25 kDa



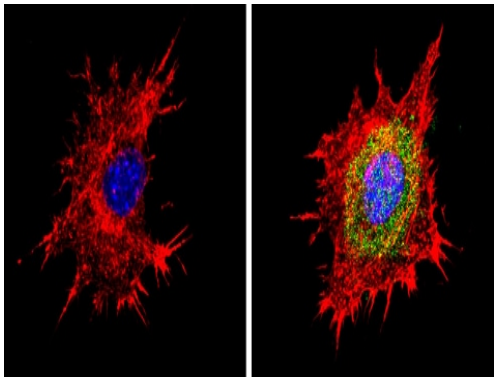
Immunocytochemistry/ Immunofluorescence - Anti-Proteasome 20S beta 6 antibody (ab3331)

Immunocytochemistry/Immunofluorescence analysis of Proteasome 20S beta 6 (green) showing staining in the cytoplasm and nucleus of A431 (Human epidermoid carcinoma cell line) 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 probed with ab3331 in 3% BSA-PBS at a dilution of 1:20 and incubated overnight at 4ºC 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.



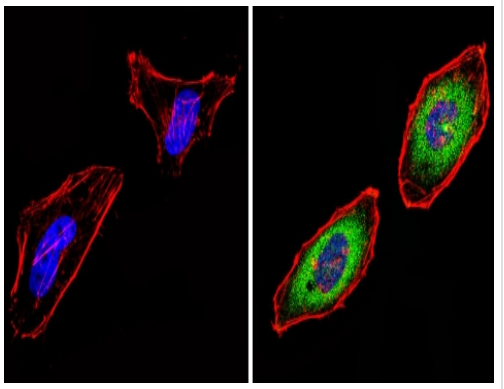
Immunocytochemistry/ Immunofluorescence - Anti-Proteasome 20S beta 6 antibody (ab3331)

Immunofluorescent analysis of Proteasome 20S Y (green) in NIH/3T3 (Mouse embryo fibroblast cell line) cells. The cells were fixed with 4% paraformaldehyde for 15 minutes, permeabilized with 0.1% Triton X-100 in PBS for 15 minutes and blocked with 3% Blocker BSA in PBS for 30 minutes at room temperature. Cells were stained with or without Proteasome 20S Y rabbit polyclonal antibody, at a concentration of 5 µg/mL for 1 hour at room temperature, and then incubated with a Goat anti-Rabbit (H+L) Superclonal Secondary Antibody, Alexa Fluor® 488 conjugate at 1/1000 dilution for 1 hour at room temperature (both panels, green). Nuclei (both panels, blue) were stained with Hoechst 33342 dye. Images were taken at 20X magnification.



Immunocytochemistry/ Immunofluorescence - Anti-Proteasome 20S beta 6 antibody (ab3331)

Immunocytochemistry/Immunofluorescence analysis of Proteasome 20S beta 6 (green) showing staining in the cytoplasm and nucleus of BAEC (Bovine aortic endothelial cell line) 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 probed with ab3331 in 3% BSA-PBS at a dilution of 1:20 and incubated overnight at 4ºC 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 20S beta 6 antibody (ab3331)

Immunocytochemistry/Immunofluorescence analysis of Proteasome 20S beta 6 (green) showing staining in the cytoplasm and nucleus of HeLa (Human epithelial adenocarcinoma cell line) 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 probed with ab3331 in 3% BSA-PBS at a dilution of 1:20 and incubated overnight at 4ºC 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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