

Anti-Cleaved Caspase-3 antibody [E83-77] ab32042

敲除验证
重组
RabMAb

★★★★☆
27 Abreviews
463 References
7 图像

概述

产品名称	Anti-Cleaved Caspase-3抗体[E83-77]
描述	兔单克隆抗体[E83-77] to Cleaved Caspase-3
宿主	Rabbit
特异性	Mouse, Rat: We have preliminary internal testing data to indicate this antibody may not react with these species. Please contact us for more information. This antibody only detects the active (cleaved) form of Caspase-3 and does not recognize the pro form of Caspase-3. PubMedID 19789217 describes the detection of human cells injected into mice. The pro-caspase-3 is cleaved only when apoptosis event occurs. So, in order to detect active Caspase-3, we strongly suggest to induce your samples into apoptotic pathway.
经测试应用	<p>适用于: WB, ICC/IF</p> <p>不适用于: Flow Cyt, IHC-P or IP</p>
种属反应性	与反应: Human, Recombinant fragment
免疫原	<p>Synthetic peptide within Human Cleaved Caspase-3 aa 1-100 (N terminal). The exact sequence is proprietary. A synthetic peptide corresponding to residues following Ser29 of human Caspase 3 (N terminus of p17 subunit).</p> <p>Database link: P42574</p>
阳性对照	WB: Wild type HAP1 + 2uM Staurosporine (ab146588) for 24 hours, whole cell lysate; Jurkat cell lysate (camptothecin treated); HeLa cell lysate (staurosporine treated). ICC/IF: Hela cells (staurosporine treated); Human Vascular endothelial cells.
常规说明	<p>This product is a recombinant monoclonal antibody, which offers several advantages including:</p> <ul style="list-style-type: none"> - High batch-to-batch consistency and reproducibility - Improved sensitivity and specificity - Long-term security of supply - Animal-free production <p>For more information see here.</p> <p>Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to RabMAb[®] patents.</p>

性能

形式 Liquid

存放说明	Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.
存储溶液	pH: 7.20 Preservative: 0.01% Sodium azide Constituents: 49% PBS, 50% Glycerol (glycerin, glycerine), 0.05% BSA
纯度	Protein A purified
克隆	单克隆
克隆编号	E83-77
同种型	IgG

应用

The Abpromise guarantee **Abpromise™** 承诺保证使用ab32042于以下的经测试应用

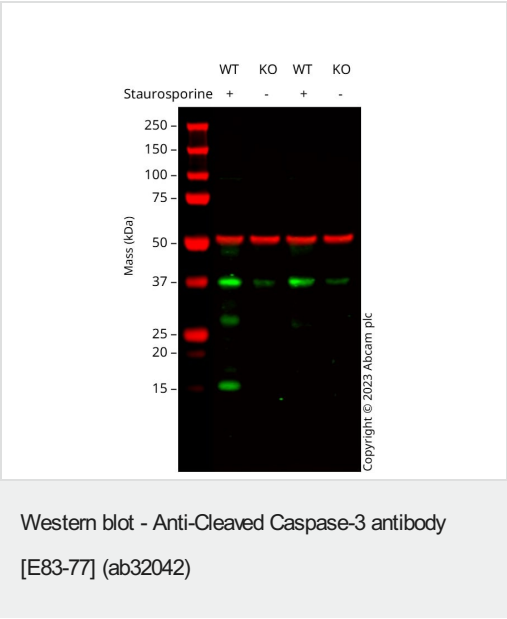
“应用说明”部分 下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
WB	★★★★★ (7)	1/500. Detects a band of approximately 17 kDa (predicted molecular weight: 32 kDa). The pro-caspase-3 is cleaved only when apoptosis event occurs. So, in order to detect active Caspase-3, we strongly suggest to induce your samples into apoptotic pathway.
ICC/IF	★★★★★ (5)	1/100 - 1/250.

应用说明 Is unsuitable for Flow Cyt, IHC-P or IP.

靶标

功能	Involved in the activation cascade of caspases responsible for apoptosis execution. At the onset of apoptosis it proteolytically cleaves poly(ADP-ribose) polymerase (PARP) at a '216-Asp-Gly-217' bond. Cleaves and activates sterol regulatory element binding proteins (SREBPs) between the basic helix-loop-helix leucine zipper domain and the membrane attachment domain. Cleaves and activates caspase-6, -7 and -9. Involved in the cleavage of huntingtin. Triggers cell adhesion in sympathetic neurons through RET cleavage.
组织特异性	Highly expressed in lung, spleen, heart, liver and kidney. Moderate levels in brain and skeletal muscle, and low in testis. Also found in many cell lines, highest expression in cells of the immune system.
序列相似性	Belongs to the peptidase C14A family.
翻译后修饰	Cleavage by granzyme B, caspase-6, caspase-8 and caspase-10 generates the two active subunits. Additional processing of the propeptides is likely due to the autocatalytic activity of the activated protease. Active heterodimers between the small subunit of caspase-7 protease and the large subunit of caspase-3 also occur and vice versa. S-nitrosylated on its catalytic site cysteine in unstimulated human cell lines and denitrosylated upon activation of the Fas apoptotic pathway, associated with an increase in intracellular caspase activity. Fas therefore activates caspase-3 not only by inducing the cleavage of the caspase zymogen to its active subunits, but also by stimulating the denitrosylation of its active site thiol.
细胞定位	Cytoplasm.



Western blot - Anti-Cleaved Caspase-3 antibody [E83-77] (ab32042)

All lanes : Anti-Cleaved Caspase-3 antibody [E83-77] (ab32042) at 1/500 dilution

Lane 1 : Wild-type HeLa Treated Staurosporine (2uM, 4h) cell lysate

Lane 2 : CASP3 knockout HeLa Treated Staurosporine (2uM, 4h) cell lysate

Lane 3 : Wild-type HeLa Vehicle Control Staurosporine (0uM, 4h) cell lysate

Lane 4 : CASP3 knockout HeLa Vehicle Control Staurosporine (0uM, 4h) cell lysate

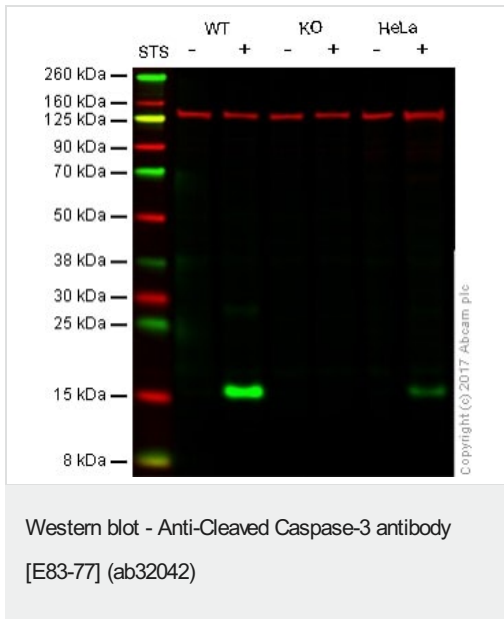
Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

Predicted band size: 32 kDa

Observed band size: 16,28 kDa

Western blot: Anti-CASP3 antibody [E83-77] (ab32042) staining at 1/500 dilution, shown in green; Mouse anti-Alpha Tubulin [DM1A] ([ab7291](#)) loading control staining at 1/20000 dilution, shown in red. In Western blot, ab32042 was shown to bind specifically to CASP3. A band was observed at 16/28 kDa in treated wild-type HeLa cell lysates with no signal observed at this size in CASP3 knockout cell line. To generate this image, wild-type and CASP3 knockout HeLa cell lysates were analysed. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in 3 % milk in TBS-0.1 % Tween[®] 20 (TBS-T) before incubation with primary antibodies overnight at 4 °C. Blots were washed four times in TBS-T, incubated with secondary antibodies for 1 h at room temperature, washed again four times then imaged. Secondary antibodies used were Goat anti-Rabbit IgG H&L 800CW and Goat anti-Mouse IgG H&L 680RD at 1/20000 dilution.



Lane 1: Wild type HAP1 + DMSO for 24 hours, whole cell lysate (20 µg)

Lane 2: Wild type HAP1 + 2uM Staurosporine ([ab146588](#)) for 24 hours, whole cell lysate (20 µg)

Lane 3: HAP1 CASP3 KO + DMSO for 24 hours, whole cell lysate (20 µg)

Lane 4: HAP1 CASP3 KO + 2uM Staurosporine ([ab146588](#)) for 24 hours, whole cell lysate (20 µg)

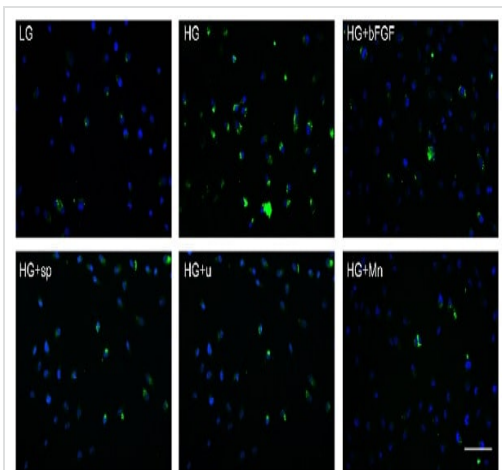
Lane 5: HeLa + DMSO for 24 hours, whole cell lysate (20 µg)

Lane 6: HeLa + 2uM Staurosporine ([ab146588](#)) for 24 hours, whole cell lysate (20 µg)

Lanes 1 - 6: Merged signal (red and green). Green - ab32042 observed at 17 kDa. Red - loading control, [ab130007](#), observed at 130 kDa.

ab32042 was shown to specifically react with CASP3 (Caspase-3) when CASP3 (Caspase-3) knockout samples were used. HAP1 wild-type and CASP3 (Caspase-3) knockout samples were subjected to SDS-PAGE. Ab32042 and [ab130007](#) (Mouse anti vinculin loading control) were incubated overnight at 4°C at 500 dilution and 1/10000 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 1/10000 dilution for 1 hour at room temperature before imaging.

Cells were grown to confluency prior to treatment.

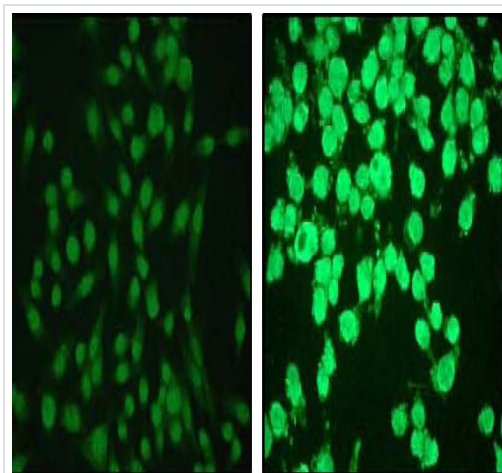


Immunocytochemistry/ Immunofluorescence - Anti-Cleaved Caspase-3 antibody [E83-77] (ab32042)

Image from Zhu ZX et al., PLoS One. 2015;10(12):e0144495. Fig 7.; doi: 10.1371/journal.pone.0144495. Reproduced under the Creative Commons license <http://creativecommons.org/licenses/by/4.0/>

High-glucose induces apoptosis in human Vascular endothelial cells (VECs).

Apoptotic responses in VEC were analyzed by detection of cleaved-Caspase-3 immunofluorescence using ab32042. Cells were treated with low glucose (LG) or high glucose (HG) for 72 hours before treated with 100 ng/mL bFGF, 1 μ M sp600125 (sp), or 1 μ M U0126 (U) or 10 μ M MnTmPyP for 1 hour. Bar = 100 μ m.

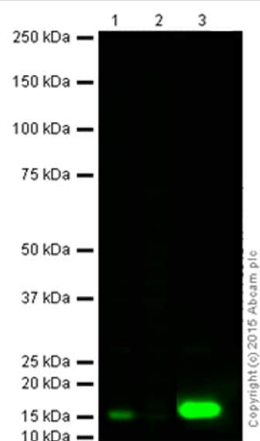


Immunocytochemistry/ Immunofluorescence - Anti-Cleaved Caspase-3 antibody [E83-77] (ab32042)

Ab32042, at dilution of 1/100, staining HeLa (human epithelial cell line from cervix adenocarcinoma) cells by Immunofluorescence.

Left image: control.

Right image: staurosporine treated.



Western blot - Anti-Cleaved Caspase-3 antibody [E83-77] (ab32042)

All lanes : Anti-Cleaved Caspase-3 antibody [E83-77] (ab32042) at 1/500 dilution

Lane 1 : HeLa Whole Cell Lysate (2 uM Staurosporine, 4Hr) at 20 µg

Lane 2 : HeLa Whole Cell Lysate (untreated) at 20 µg

Lane 3 : Cleaved Caspase 3 (recombinant protein) at 0.1 µg

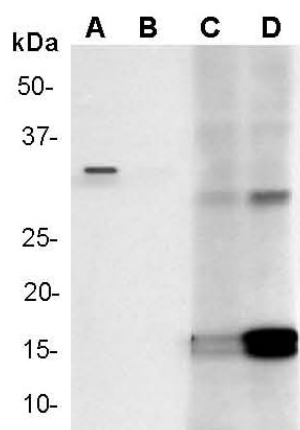
Secondary

All lanes : 800CW Goat Anti-Rabbit IgG at 1/10000 dilution

Performed under reducing conditions.

Predicted band size: 32 kDa

Additional bands at: 17 kDa (possible mature (processed) protein)



Western blot - Anti-Cleaved Caspase-3 antibody [E83-77] (ab32042)

Lane 1 : anti Pro Caspase 3 at 1/10000 dilution

Lane 2 : anti Pro Caspase 3 at 1/10000 dilution

Lanes 3-4 : Anti-Cleaved Caspase-3 antibody [E83-77] (ab32042) at 1/500 dilution

Lane 1 : Jurkat (human T cell leukemia cell line from peripheral blood) cell lysate

Lanes 2 & 4 : Jurkat cell lysate + Camptothecin

Lane 3 : Jurkat cell lysate

Predicted band size: 32 kDa

Observed band size: 17 kDa

Additional bands at: 30 kDa. We are unsure as to the identity of these extra bands.

Why choose a recombinant antibody?



Research with confidence
Consistent and reproducible results



Long-term and scalable supply
Recombinant technology



Success from the first experiment
Confirmed specificity



Ethical standards compliant
Animal-free production

Anti-Cleaved Caspase-3 antibody [E83-77]
(ab32042)

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