abcam

Product datasheet

Anti-Pro Caspase-8 antibody [EPR162] ab108333





重组 RabMAb

★★★★★ 1 Abreviews 28 References 8 图像

概述

产品名称 Anti-Pro Caspase-8抗体[EPR162]

描述 兔单克隆抗体[EPR162] to Pro Caspase-8

宿主 Rabbit

特异性 Theoretically, ab108333 should recognize p55/54, p43/41 and p18. However, in our internal tests,

it only recognized pro caspase 8. We tested it side by side with ab32397. Both antibodies are

KO validated, and ab32397 recognizes pro-caspase 8 and cleavage caspase 8.

经测试应用 适用于: WB, IHC-P

不适用于: Flow Cyt or IP

种属反应性 与反应: Mouse, Rat, Human

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

阳性对照 Jurkat, IM9, HepG2, HeLa, and HL60 cell lysates; Human tonsil tissue and Human hepatocellular

carcinoma tissue; HeLa cells.

常规说明 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**.

性能

形式 Liquid

存放说明 Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.

存储溶液 pH: 7.20

Preservative: 0.05% Sodium azide

Constituents: 0.1% BSA, 40% Glycerol (glycerin, glycerine), 9.85% Tris glycine, 50% Tissue

culture supernatant

纯度 Protein A purified

克隆 单克隆

克隆编号 EPR162

同种型 IgG

应用

The Abpromise guarantee Abpromise™承诺保证使用ab108333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		1/1000 - 1/10000. Detects a band of approximately 55 kDa (predicted molecular weight: 55 kDa).
IHC-P		1/100 - 1/250. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol. Antigen retrieval is recommended.

应用说明

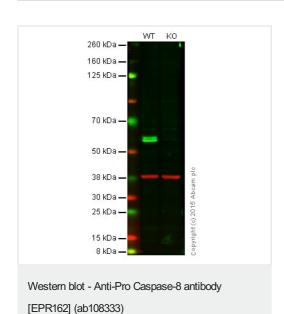
Is unsuitable for Flow Cyt or IP.

靶标

相关性

Disease: Defects in CASP8 are the cause of caspase-8 deficiency (CASP8D) [MIM:607271]. CASP8D is a disorder resembling autoimmune lymphoproliferative syndrome (ALPS). It is characterized by lymphadenopathy, splenomegaly, and defective CD95-induced apoptosis of peripheral blood lymphocytes (PBLs). It leads to defects in activation of T-lymphocytes, B-lymphocytes, and natural killer cells leading to immunodeficiency characterized by recurrent sinopulmonary and herpes simplex virus infections and poor responses to immunization.

图片

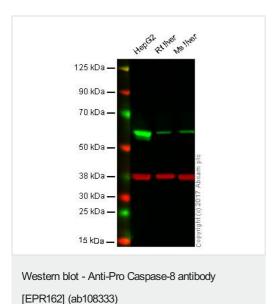


Lane 1: Wild-type HAP1 cell lysate (20 µg)

Lane 2: Pro Caspase-8 knockout HAP1 cell lysate (20 µg)

Lanes 1 and 2: Merged signal (red and green). Green - ab108333 observed at 55 kDa. Red - loading control, <u>ab8245</u>, observed at 37 kDa.

ab108333 was shown to specifically react with Caspase-8 when Caspase-8 knockout samples were used. Wild-type and Pro Caspase-8 knockout samples were subjected to SDS-PAGE. ab108333 and ab8245 (loading control to GAPDH) were both diluted 1/1000 and incubated overnight at 4°C. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1/10 000 dilution for 1 h at room temperature before imaging.



All lanes : Anti-Pro Caspase-8 antibody [EPR162] (ab108333) at 1/1000 dilution

Lane 1 : HepG2 whole cell lysate

Lane 2: Rat liver tissue lysate

Lane 3: Mouse liver tissue lysate

Lysates/proteins at 20 µg per lane.

Secondary

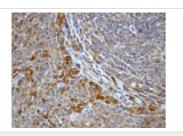
All lanes : Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (**ab216773**) at 1/10000 dilution

Performed under reducing conditions.

Predicted band size: 55 kDa **Observed band size:** 55 kDa

Lanes 1 - 3: Merged signal (red and green). Green - ab108333 observed at 55 kDa. Red - loading control, **ab8245**, observed at 37 kDa.

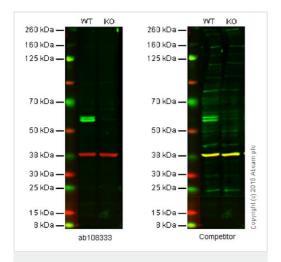
This blot was produced using a 4-12% Bis-tris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using Licor blocking buffer before being incubated with ab108333 and ab8245 (loading control) overnight at 4°C. Antibody binding was detected using Goat anti-Rabbit IgG H&L (IRDye® 800CW) preabsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed (ab216776) at a 1:10000 dilution for 1hr at room temperature and then imaged.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Pro Caspase-8 antibody [EPR162] (ab108333)

ab108333, at 1/100 dilution, staining Pro Caspase-8 in paraffinembedded human tonsil tissue by Immunohistochemistry.

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Western blot - Anti-Pro Caspase-8 antibody [EPR162] (ab108333)

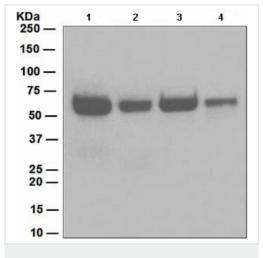


Lane 2: Pro Caspase-8 knockout HAP1 cell lysate (20 µg)

Lanes 1 and 2: Merged signal (red and green).

Green - target observed at 55 kDa. Red - loading control, <u>ab8245</u>, observed at 37 kDa.

This western blot image is a comparison between ab108333 and a competitor's top cited mouse monoclonal antibody.



Western blot - Anti-Pro Caspase-8 antibody [EPR162] (ab108333)

All lanes : Anti-Pro Caspase-8 antibody [EPR162] (ab108333) at 1/1000 dilution

Lane 1 : Jurkat cell lysate

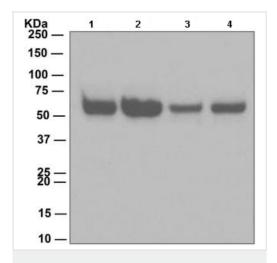
Lane 2: IM9 cell lysate

Lane 3: HepG2 cell lysate

Lane 4: HL60 cell lysate

Lysates/proteins at 10 µg per lane.

Predicted band size: 55 kDa



Western blot - Anti-Pro Caspase-8 antibody [EPR162] (ab108333)

All lanes : Anti-Pro Caspase-8 antibody [EPR162] (ab108333) at 1/1000 dilution

Lane 1: Jurkat cell lysate, treated with etoposide

Lane 2: Jurkat cell lysate

Lane 3: HeLa cell lysate, treated staurosporine

Lane 4: HeLa cell lysate

Lysates/proteins at 10 µg per lane.

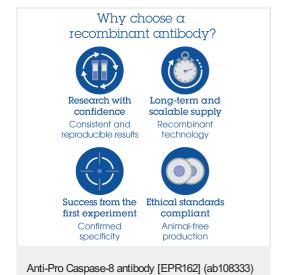
Predicted band size: 55 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Pro Caspase-8 antibody [EPR162] (ab108333)

ab108333, at 1/100 dilution, staining Pro Caspase-8 in paraffinembedded human hepatocellular carcinoma by Immunohistochemistry.

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



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