# abcam

#### Product datasheet

## Anti-CD8 alpha antibody [OX-8] ab33786

★★★★★ 4 Abreviews 33 References 5 图像

#### 概述

产品名称 Anti-CD8 alpha抗体[OX-8]

**小鼠单克隆抗体[OX-8] to CD8 alpha** 

**宿主** Mouse

经测试应用 适用于: Flow Cyt, ICC/IF, IHC-P, WB

种属反应性 **与反**应: Rat

免疫原 Tissue, cells or virus. High molecular weight rat thymocyte glycoproteins

阳性对照 WB: Rat thymus tissue lysate. IHC-P: Rat Spleen. Flow Cyt: Rat splenocytes. ICC/IF: Rat

splenocyte cells.

常规说明

This antibody clone is manufactured by Abcam. If you require a custom buffer formulation or

conjugation for your experiments, please contact orders@abcam.com.

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 long

term. Avoid freeze / thaw cycle.

**存储溶液** pH: 7.20

Preservative: 0.02% Sodium azide Constituents: PBS, 6.97% L-Arginine

纯**度** Protein G purified

 克隆
 单克隆

 克隆编号
 OX-8

 骨髓瘤
 NS1

1

同种型 lgG1

轻链类型 kappa

#### 应用

### The Abpromise guarantee Abpromise™承诺保证使用ab33786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。

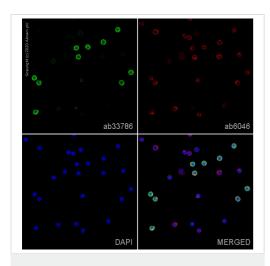
应用	Ab评论	说明
Flow Cyt		1/125000.
ICC/IF		Use at an assay dependent concentration.
IHC-P	<b>★★★★★ (2)</b>	Use a concentration of 5 µg/ml.
WB		Use a concentration of 1 µg/ml. Predicted molecular weight: 26 kDa.

功能	Identifies cytotoxic/suppressor T-cells that interact with MHC class I bearing targets. CD8 is thought to play a role in the process of T-cell mediated killing. CD8 alpha chains binds to class I MHC molecules alpha-3 domains.
疾病相关	Defects in CD8A are a cause of familial CD8 deficiency (CD8 deficiency) [MIM:608957]. Familial CD8 deficiency is a novel autosomal recessive immunologic defect characterized by absence of CD8+ cells, leading to recurrent bacterial infections.
序列相似性	Contains 1 lg-like V-type (immunoglobulin-like) domain.
翻译后修饰	All of the five most carboxyl-terminal cysteines form inter-chain disulfide bonds in dimers and higher multimers, while the four N-terminal cysteines do not.

细**胞定位** Secreted and Cell membrane.

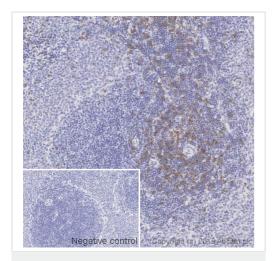
#### 图片

靶标



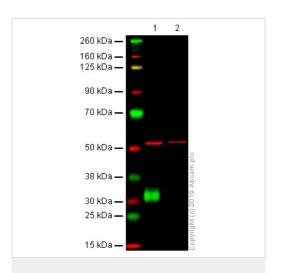
Immunocytochemistry/ Immunofluorescence - Anti-CD8 alpha antibody [OX-8] (ab33786)

ab33786 staining CD8 alpha in Rat splenocytes cells. The cells were fixed with 4% paraformaldehyde (10 min), permeabilized with 0.1% PBS-Tween for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated overnight at 4°C with ab33786 at 5µg/ml and ab6046, Rabbit polyclonal to beta Tubulin - Loading Control. Cells were then incubated with ab150117, Goat polyclonal Secondary Antibody to Mouse IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 dilution (shown in green) and ab150080, Goat polyclonal Secondary Antibody to Rabbit IgG - H&L (Alexa Fluor® 594) at 1/1000 dilution (shown in pseudocolour red). Nuclear DNA was labelled with DAPI (shown in blue). Also suitable in cells fixed with 100% methanol (5 min).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD8 alpha antibody [OX-8] (ab33786)

IHC image of CD8 alpha staining in a section of formalin-fixed paraffin-embedded normal Rat Spleen performed on a Leica BOND™ system using the standard Protocol F. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20mins. The section was then incubated with <a href="mailto:ab33923">ab33923</a>, 5µg/ml, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX. The inset secondary-only control image is taken from an identical assay without primary antibody. For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.



Western blot - Anti-CD8 alpha antibody [OX-8] (ab33786)

All lanes : Anti-CD8 alpha antibody [OX-8] (ab33786) at 1  $\mu$ g/ml

Lane 1 : Rat thymus tissue lysate

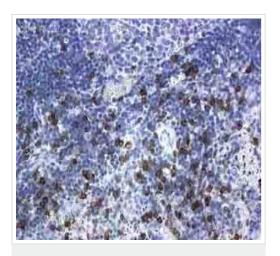
Lane 2 : Rat brain tissue lysate

Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

**Predicted band size:** 26 kDa **Observed band size:** 32 kDa

This blot was produced using a 4-12% Bis-tris under the MOPS buffer system. The gel was run at 200V for 55 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was blocked for an hour using 3% milk before ab33786 and <a href="mailto:ab176560">ab176560</a> (Rabbit anti-alpha Tubulin loading control) were incubated overnight at 4°C at a 1 µg/ml concentration and 1/20000 dilution respectively. Antibody binding was detected using Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse lgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1/20000 dilution for 1 hour at room temperature before imaging.

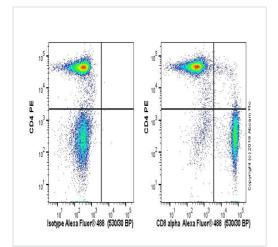


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD8 alpha antibody [OX-8] (ab33786)

Image courtesy of an anonymous Abreview.

ab33786 staining CD8 alpha in rat spleen tissue by Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections).

Tissue was fixed with 10% neutral buffered formalin and a heat mediated antigen retrieval step was performed using citrate buffer. Samples were then permeabilized with Triton X-100, blocked using 5% serum for 30 minutes at 25°C and then incubated with ab33786 at a 1/100 dilution for 1 hour at 25°C. The secondary used was a biotin-conjugated rabbit anti-mouse polyclonal, used at a 1/300 dilution.



Flow Cytometry - Anti-CD8 alpha antibody [OX-8] (ab33786)

Flow cytometry staining of Lewis rat splenocytes with ab33786 (right) or mouse lgG1 kappa; (ab170190) isotype (left). Cells were incubated for 30 min on ice in 1x PBS containing 10 % rat serum to block FC receptors and non-specific protein-protein interaction followed by the antibody (ab33786) or mouse lgG1 kappa; (ab170190) isotype (1x  $10^6$  in 100  $\mu$ l at 0.008  $\mu$ g/ml) for 30 min on ice.

The secondary antibody Goat anti-mouse IgG H&L (Alexa Fluor  $^{@}$  488, pre-adsorbed) (<u>ab150177</u>) was used at 1/2000 dilution for 30 min on ice.

The cells were simultaneously stained with CD4.

Acquisition of >30,000 events were collected using a 50 mW Blue laser (488nm) and 530/30 bandpass filter. Events were gated on live CD3 positive T cells.

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