


Anti-alpha 2 Macroglobulin antibody ab58703

★★★★★ [5 Abreviews](#) [10 References](#) [3 图像](#)

概述

产品名称	Anti-alpha 2 Macroglobulin抗体
描述	兔多克隆抗体to alpha 2 Macroglobulin
宿主	Rabbit
经测试应用	适用于: WB, IHC-P
种属反应性	与反应: Human 预测可用于: Mouse 
免疫原	Synthetic peptide corresponding to C terminal residues of human alpha 2 Macroglobulin.
常规说明	<p>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.</p> <p>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</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: 7.2 Preservative: 0.01% Sodium azide Constituents: PBS, 50% Glycerol (glycerin, glycerine)
纯度	Immunogen affinity purified
纯化说明	Purified by antigen specific affinity chromatography.
克隆	多克隆
同种型	IgG

应用

The Abpromise guarantee

Abpromise™ 承诺保证使用ab58703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。

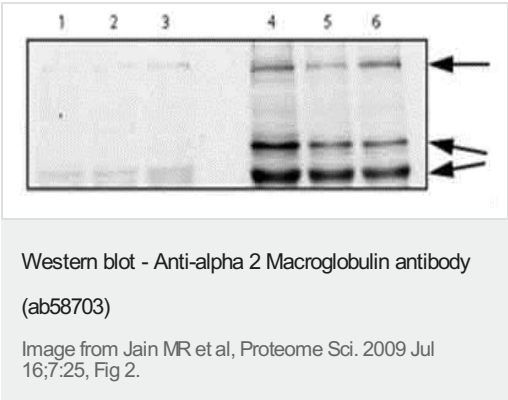
应用	Ab评论	说明
WB	★★★★★ (2)	
IHC-P	★★★★★ (2)	

应用说明	<p>ELISA: Use at an assay dependent dilution.</p> <p>ICC: 1/500.</p> <p>IHC-P: Use at a concentration of 2 µg/ml.</p> <p>WB: Use at an assay dependent concentration. Predicted molecular weight: 163 kDa.</p> <p>Not yet tested in other applications.</p> <p>Optimal dilutions/concentrations should be determined by the end user.</p>
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靶标

功能	<p>Is able to inhibit all four classes of proteinases by a unique 'trapping' mechanism. This protein has a peptide stretch, called the 'bait region' which contains specific cleavage sites for different proteinases. When a proteinase cleaves the bait region, a conformational change is induced in the protein which traps the proteinase. The entrapped enzyme remains active against low molecular weight substrates (activity against high molecular weight substrates is greatly reduced). Following cleavage in the bait region a thioester bond is hydrolyzed and mediates the covalent binding of the protein to the proteinase.</p>
组织特异性	<p>Secreted in plasma.</p>
序列相似性	<p>Belongs to the protease inhibitor I39 (alpha-2-macroglobulin) family.</p>
发展阶段	<p>Contrary to the rat protein, which is an acute phase protein, this protein is always present at high levels in circulation.</p>
细胞定位	<p>Secreted.</p>

图片



All lanes : Anti-alpha 2 Macroglobulin antibody (ab58703) at 1/1000 dilution

Lanes 1-3 : lysates prepared from rat spinal cord tissues (control group)

Lanes 4-6 : lysates prepared from rat spinal cord tissues (encephalomyelitis group)

Lysates/proteins at 30 µg per lane.

Secondary

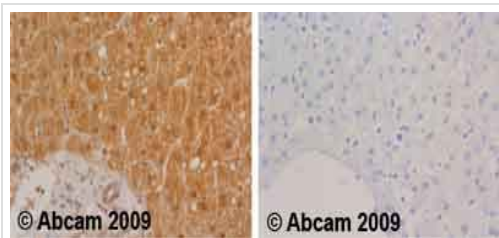
All lanes : goat anti-rabbit IgG coupled to horseradish peroxidase

Predicted band size: 163 kDa

Observed band size: 163 kDa

Additional bands at: 60 kDa (possible cleavage fragment), 80 kDa (possible cleavage fragment)

Three independent control and encephalomyelitis samples were used for Western blotting analysis. Samples were first resolved on a 12.5% SDS-PAGE gel and then transferred to a nitrocellulose membrane. The membrane was rinsed with PBS and the non specific binding sites were blocked in a solution of 5% non-fat milk in PBST (0.05% Tween 20 in PBS) for 1 hour at room temperature, followed by three washes in PBST for 10 minutes each. The membrane was first incubated with ab58703 at a 1/1,000 dilution overnight and then washed in PBST buffer as described above. The immunocomplexes were visualized by Western Lightning® Western Blot Chemiluminescence Reagent Plus, using a goat anti-rabbit IgG coupled to horseradish peroxidase as the secondary antibody.

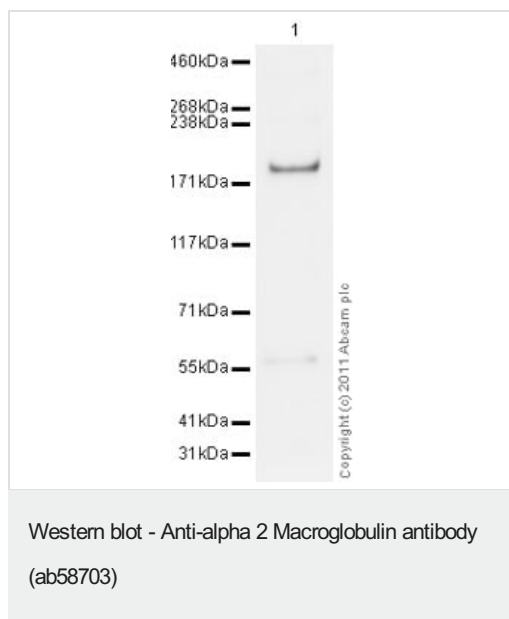


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-alpha 2 Macroglobulin antibody (ab58703)

Ab58703 staining alpha 2 Macroglobulin in human liver.

Left panel: with primary antibody at 2 ug/ml. Right panel: isotype control.

Sections were stained using an automated system DAKO Autostainer Plus , at room temperature. Sections were rehydrated and antigen retrieved with the Dako 3-in-1 antigen retrieval buffers EDTA pH 9.0 in a DAKO PT Link. Slides were peroxidase blocked in 3% H₂O₂ in methanol for 10 minutes. They were then blocked with Dako Protein block for 10 minutes (containing casein 0.25% in PBS) then incubated with primary antibody for 20 minutes and detected with Dako Envision Flex amplification kit for 30 minutes. Colorimetric detection was completed with diaminobenzidine for 5 minutes. Slides were counterstained with Haematoxylin and coverslipped under DePeX. Please note that for manual staining we recommend to optimize the primary antibody concentration and incubation time (overnight incubation), and amplification may be required.



Anti-alpha 2 Macroglobulin antibody (ab58703) at 1 µg/ml + Native Human alpha 2 Macroglobulin protein ([ab77935](#)) at 0.1 µg

Secondary

Goat Anti-Rabbit IgG H&L (HRP) preadsorbed ([ab97080](#)) at 1/5000 dilution

Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 163 kDa

Exposure time: 3 minutes

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