

Anti-Endothelin 1 antibody [TR.ET.48.5] ab2786

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概述

产品名称	Anti-Endothelin 1 抗体[TR.ET.48.5]
描述	小鼠单克隆抗体[TR.ET.48.5] to Endothelin 1
宿主	Mouse
特异性	Immunohistochemical staining of ET-1 in human corpus cavernosum tissue with this antibody results in staining of endothelial cells. Radioimmune assays can be used to concentrate ET-1 in solution (e.g. serum/plasma, milk, urine).
经测试应用	适用于: Flow Cyt, ICC/IF, IHC-P, WB
种属反应性	与反应: Mouse, Rat, Human
免疫原	Full length native protein (purified) corresponding to Human Endothelin 1 conjugated to keyhole limpet haemocyanin.
表位	Studies suggest that this antibody binds to an epitope in the region of ET-1 represented by amino acids 8-16.
阳性对照	human bowel
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. 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	Preservative: 0.05% Sodium azide Constituent: PBS
纯度	Protein G purified
克隆	单克隆

克隆编号 TR.ET.48.5
同种型 IgG1

应用

The Abpromise guarantee **Abpromise™** 承诺保证使用 ab2786 于以下的经测试应用

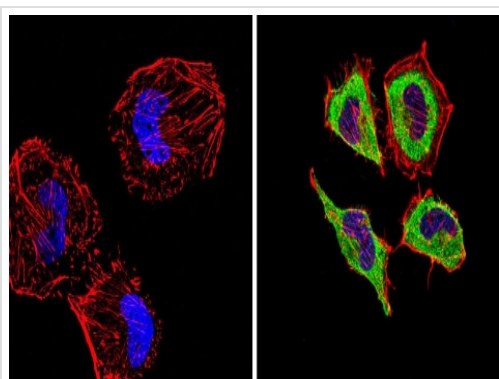
“应用说明”部分 下显示的仅为推荐的起始稀释度；实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
Flow Cyt		Use at an assay dependent concentration. ab170190 - Mouse monoclonal IgG1, is suitable for use as an isotype control with this antibody.
ICC/IF		1/200 - 1/1000.
IHC-P		1/250.
WB		Use at an assay dependent concentration. Predicted molecular weight: 24 kDa.

靶标

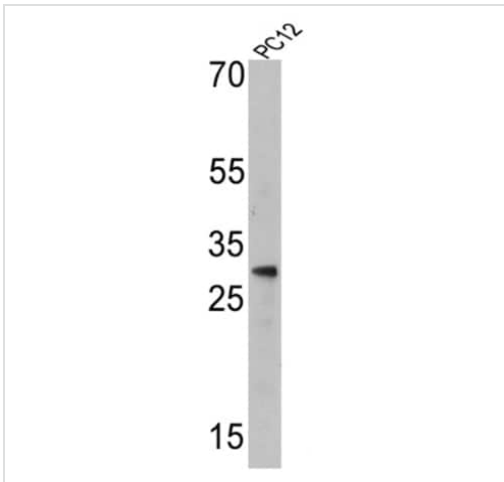
功能 Endothelins are endothelium-derived vasoconstrictor peptides.
组织特异性 Expressed in lung, placental stem villi vessels and in cultured placental vascular smooth muscle cells.
序列相似性 Belongs to the endothelin/sarafotoxin family.
细胞定位 Secreted.

图片



Immunocytochemistry/ Immunofluorescence - Anti-Endothelin 1 antibody [TR.ET.48.5] (ab2786)

ab2786 labelling Endothelin 1 (green) in the secretion of HeLa cells (right) compared with a negative control (left) by Immunocytochemistry/Immunofluorescence. 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 incubated with the primary antibody (1:200 in 3% BSA-PBS) overnight at 4 °C. A DyLight 488-conjugated Goat anti-mouse IgG (H+L) was used as the secondary antibody. Red (phalloidin) - F-actin, Blue (DAPI) - nuclei. Images were taken at a magnification of 60x.

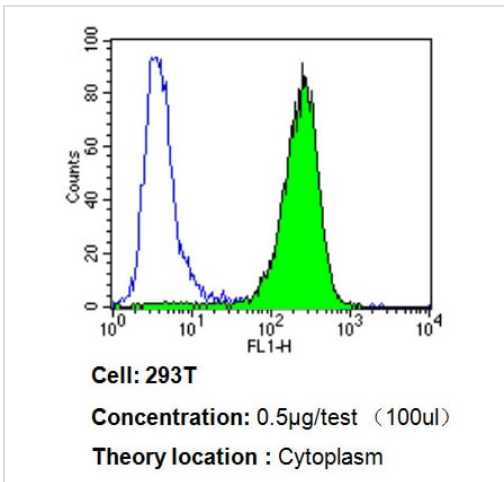


Western blot - Anti-Endothelin 1 antibody
[TR.ET.48.5] (ab2786)

Anti-Endothelin 1 antibody [TR.ET.48.5] (ab2786) at 1/500 dilution
+ PC12 cell lysate at 25 µg

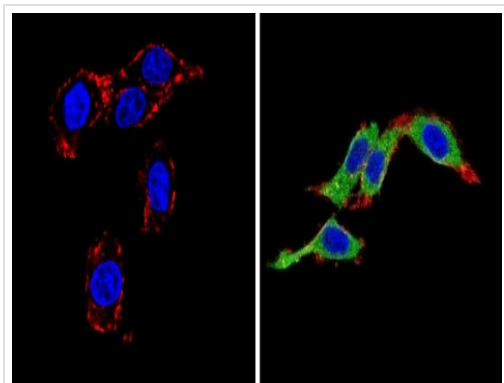
Predicted band size: 24 kDa

Observed band size: 30 kDa



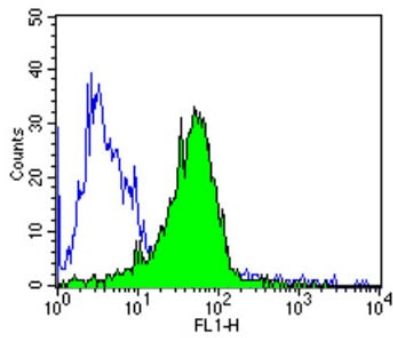
Flow Cytometry - Anti-Endothelin 1 antibody
[TR.ET.48.5] (ab2786)

Flow cytometry analysis of Endothelin 1 showing positive staining in the cytoplasm of 293T cells compared to an isotype control (blue). Cells were harvested, adjusted to a concentration of $1-5 \times 10^6$ cells/ml, fixed with 2% paraformaldehyde, washed with PBS, and incubated with ab2786 (0.5 µg/test) for 60 min at room temperature. Cells were then blocked in a solution of 2% BSA-PBS for 30 min at room temperature, incubated for 40 min at room temperature in the dark using a Dylight 488-conjugated goat anti-mouse IgG (H+L) secondary antibody, and re-suspended in PBS for FACS analysis.



Immunocytochemistry/ Immunofluorescence - Anti-Endothelin 1 antibody [TR.ET.48.5] (ab2786)

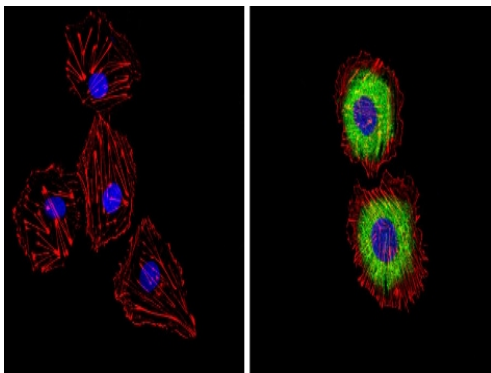
ab2786 labelling Endothelin 1 (green) in the secretion of PC12 cells (right) compared with a negative control (left) by Immunocytochemistry/Immunofluorescence. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were incubated with the primary antibody (1:200 in 3% BSA-PBS) overnight at 4 °C. A DyLight 488-conjugated Goat anti-mouse IgG (H+L) was used as the secondary antibody. Red (phalloidin) - F-actin, Blue (DAPI) - nuclei. Images were taken at a magnification of 60x.



Cell: HepG2
Concentration: 0.5µg/test (100ul)
Theory location : Cytoplasm

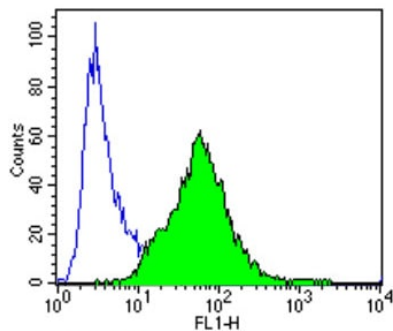
Flow Cytometry - Anti-Endothelin 1 antibody
 [TR.ET.48.5] (ab2786)

Flow cytometry analysis of Endothelin 1 showing positive staining in the cytoplasm of HepG2 cells compared to an isotype control (blue). Cells were harvested, adjusted to a concentration of $1-5 \times 10^6$ cells/ml, fixed with 2% paraformaldehyde, washed with PBS, and incubated with ab2786 (0.5 µg/test) for 60 min at room temperature. Cells were then blocked in a solution of 2% BSA-PBS for 30 min at room temperature, incubated for 40 min at room temperature in the dark using a Dylight 488-conjugated goat anti-mouse IgG (H+L) secondary antibody, and re-suspended in PBS for FACS analysis.



Immunocytochemistry/ Immunofluorescence - Anti-Endothelin 1 antibody [TR.ET.48.5] (ab2786)

ab2786 labelling Endothelin 1 (green) in the secretion of HUVEC cells (right) compared with a negative control (left) by Immunocytochemistry/Immunofluorescence. 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 incubated with the primary antibody (1:200 in 3% BSA-PBS) overnight at 4 °C. A Dylight 488-conjugated Goat anti-mouse IgG (H+L) was used as the secondary antibody. Red (phalloidin) - F-actin, Blue (DAPI) - nuclei. Images were taken at a magnification of 60x.



Cell: 3T3
Concentration: 0.5µg/test (100ul)
Theory location : Cytoplasm

Flow Cytometry - Anti-Endothelin 1 antibody
 [TR.ET.48.5] (ab2786)

Flow cytometry analysis of Endothelin 1 showing positive staining in the cytoplasm of 3T3 cells compared to an isotype control (blue). Cells were harvested, adjusted to a concentration of $1-5 \times 10^6$ cells/ml, fixed with 2% paraformaldehyde, washed with PBS, and incubated with ab2786 (0.5 µg/test) for 60 min at room temperature. Cells were then blocked in a solution of 2% BSA-PBS for 30 min at room temperature, incubated for 40 min at room temperature in the dark using a Dylight 488-conjugated goat anti-mouse IgG (H+L) secondary antibody, and re-suspended in PBS for FACS analysis.

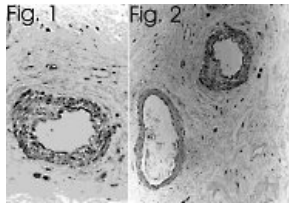


Figure 1 and Figure 2 show immunolocalization of ET-1 in human bowel using ab2786.

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Endothelin 1 antibody [TR.ET.48.5] (ab2786)

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