

## Product datasheet

# Anti-Smad2 antibody [EP784Y] ab40855

RabMAb

★★★★★ 4 Abreviews 7 References 10 图像

### 概述

<b>产品名称</b>	Anti-Smad2抗体[EP784Y]
<b>描述</b>	兔单克隆抗体[EP784Y] to Smad2
<b>特异性</b>	This antibody is specific for MH 1 domain of Smad2.
<b>经测试应用</b>	<b>适用于:</b> IHC-P, IP, ICC/IF, WB, Flow Cyt
<b>种属反应性</b>	<b>与反应:</b> Rat, Human
<b>免疫原</b>	Synthetic peptide (the amino acid sequence is considered to be commercially sensitive)
<b>阳性对照</b>	Jurkat cell lysate and human prostate carcinoma tissue.
<b>常规说明</b>	<p>This product is a recombinant rabbit monoclonal antibody.</p> <p><b>The rat recommendation is based on the WB results. This antibody may not be suitable for IHC with rat samples.</b></p> <p>Mouse: We have preliminary internal testing data to indicate this antibody may not react with these species. Please contact us for more information.</p> <p>Our RabMAb<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <a href="#">RabMAb<sup>®</sup> patents</a></p> <p><b>We are constantly working hard to ensure we provide our customers with best in class antibodies. As a result of this work we are pleased to now offer this antibody in purified format. We are in the process of updating our datasheets. The purified format is designated 'PUR' on our product labels. If you have any questions regarding this update, please contact our Scientific Support team.</b></p>

### 性能

<b>形式</b>	Liquid
<b>存放说明</b>	Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.
<b>存储溶液</b>	<p>pH: 7.20</p> <p>Preservative: 0.01% Sodium azide</p> <p>Constituents: 40% Glycerol, 0.05% BSA, 59% PBS</p>
<b>纯度</b>	Protein A purified
<b>克隆</b>	单克隆

克隆编号	EP784Y
同种型	IgG

## 应用

Our [Abpromise guarantee](#) covers the use of **ab40855** in the following tested applications.

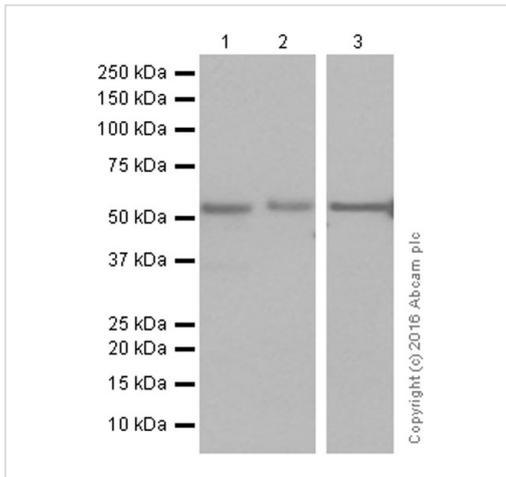
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

应用	Ab评论	说明
IHC-P		1/50.
IP		1/20 - 1/50.
ICC/IF		1/100 - 1/250.
WB	★★★★★	1/2000 - 1/10000. Detects a band of approximately 55 kDa (predicted molecular weight: 58 kDa).
Flow Cyt		1/20 - 1/100. <a href="#">ab172730</a> - Rabbit monoclonal IgG, is suitable for use as an isotype control with this antibody.

## 靶标

功能	Receptor-regulated SMAD (R-SMAD) that is an intracellular signal transducer and transcriptional modulator activated by TGF-beta (transforming growth factor) and activin type 1 receptor kinases. Binds the TRE element in the promoter region of many genes that are regulated by TGF-beta and, on formation of the SMAD2/SMAD4 complex, activates transcription. May act as a tumor suppressor in colorectal carcinoma.
组织特异性	Expressed at high levels in skeletal muscle, heart and placenta.
序列相似性	Belongs to the dwarfin/SMAD family. Contains 1 MH1 (MAD homology 1) domain. Contains 1 MH2 (MAD homology 2) domain.
翻译后修饰	Phosphorylated on one or several of Thr-220, Ser-245, Ser-250, and Ser-255. In response to TGF-beta, phosphorylated on Ser-465/467 by TGF-beta and activin type 1 receptor kinases. Able to interact with SMURF2 when phosphorylated on Ser-465/467, recruiting other proteins, such as SNON, for degradation. In response to decorin, the naturally occurring inhibitor of TGF-beta signaling, phosphorylated on Ser-240 by CaMK2. Phosphorylated by MAPK3 upon EGF stimulation; which increases transcriptional activity and stability, and is blocked by calmodulin. In response to TGF-beta, ubiquitinated by NEDD4L; which promotes its degradation. Acetylated on Lys-19 by coactivators in response to TGF-beta signaling, which increases transcriptional activity. Isoform short: Acetylation increases DNA binding activity in vitro and enhances its association with target promoters in vivo. Acetylation in the nucleus by EP300 is enhanced by TGF-beta.
细胞定位	Cytoplasm. Nucleus. Cytoplasmic and nuclear in the absence of TGF-beta. On TGF-beta stimulation, migrates to the nucleus when complexed with SMAD4. On dephosphorylation by phosphatase PPM1A, released from the SMAD2/SMAD4 complex, and exported out of the nucleus by interaction with RANBP1.

## 图片



Western blot - Anti-Smad2 antibody [EP784Y] (ab40855)

**All lanes** : Anti-Smad2 antibody [EP784Y] (ab40855) at 1/2000 dilution

**Lane 1** : A-673 (Human muscle Ewing's Sarcoma cell line) whole cell lysate

**Lane 2** : HUVEC (Human umbilical vein endothelial cell line) whole cell lysate

**Lane 3** : C6 (Rat glial tumor cell line) whole cell lysate

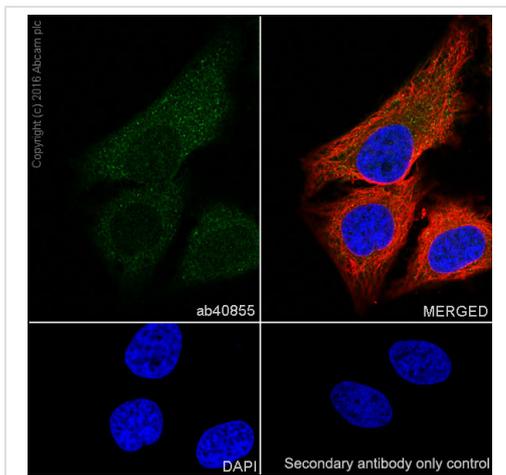
Lysates/proteins at 20 µg per lane.

### Secondary

Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution (Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated)

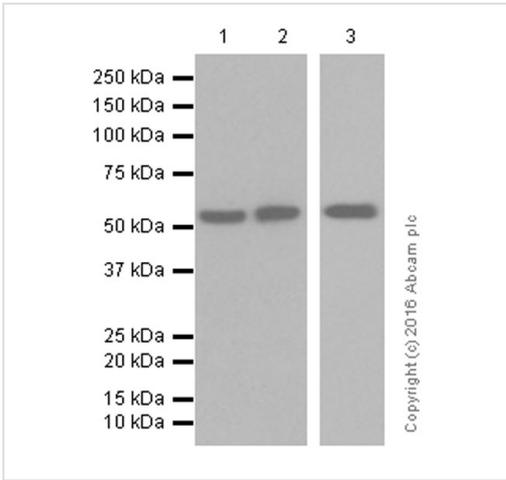
**Predicted band size** : 58 kDa

Diluting and blocking buffer: 5% NFDM /TBST



Immunocytochemistry/ Immunofluorescence - Anti-Smad2 antibody [EP784Y] (ab40855)

ab40855 staining Smad2 in HeLa (human cervix adenocarcinoma) cells by ICC/IF (Immunocytochemistry/immunofluorescence). Cells were fixed with 4% Paraformaldehyde and permeabilized with 0.1% Triton X-100. Samples were incubated with primary antibody at a dilution of 1/100. A goat anti rabbit IgG (Alexa Fluor® 488) (ab150077) was used as the secondary antibody at a dilution of 1/1000. ab195889 was used as a counterstain for primary antibody ab40855 at 1/1000. DAPI was used as a nuclear counterstain and PBS as a negative control.



Western blot - Anti-Smad2 antibody [EP784Y] (ab40855)

**All lanes :** Anti-Smad2 antibody [EP784Y] (ab40855) at 1/10000 dilution

**Lane 1 :** Jurkat (Human T cell leukemia cell line from peripheral blood) whole cell lysate

**Lane 2 :** HeLa (Human epithelial cell line from cervix adenocarcinoma) whole cell lysate

**Lane 3 :** HeLa (Human epithelial cell line from cervix adenocarcinoma) whole cell lysate

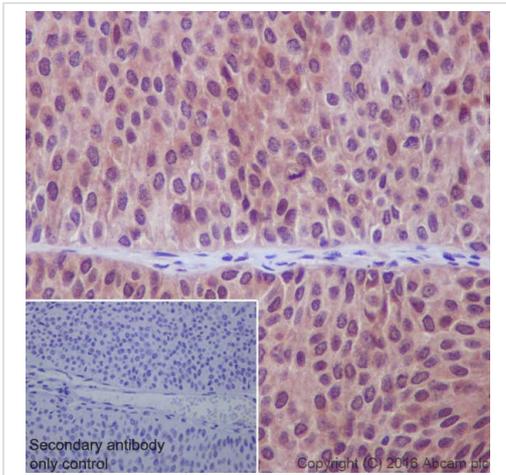
Lysates/proteins at 20 µg per lane.

**Secondary**

Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 20000 µg (Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated)

**Predicted band size :** 58 kDa

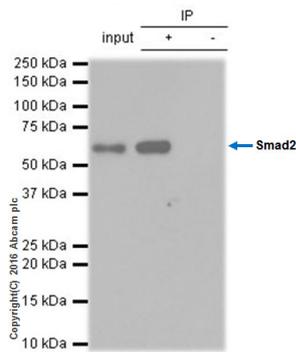
Blocking and diluting buffer: 5% NFDm/TBST



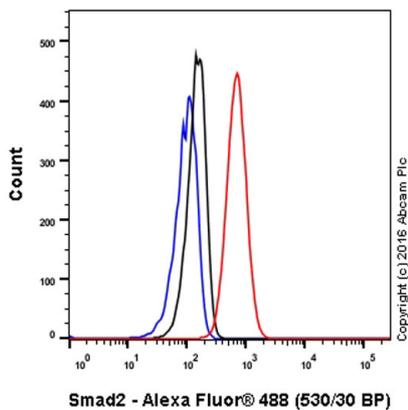
Immunohistochemistry - Anti-Smad2 antibody [EP784Y] (ab40855)

ab40855 staining Smad2 in human bladder carcinoma tissue sections by Immunohistochemistry (IHC-P - paraformaldehyde-fixed, paraffin-embedded sections). Tissue was fixed with paraformaldehyde and antigen retrieval was by heat mediation in a EDTA buffer. Samples were incubated with primary antibody at a dilution of 1/50. A ImmunoHistoProbe one step HRP Polymer was used as a secondary antibody, ready to use.

**Negative control 1:** PBS in place of primary antibody.



Immunoprecipitation - Anti-Smad2 antibody  
[EP784Y] (ab40855)

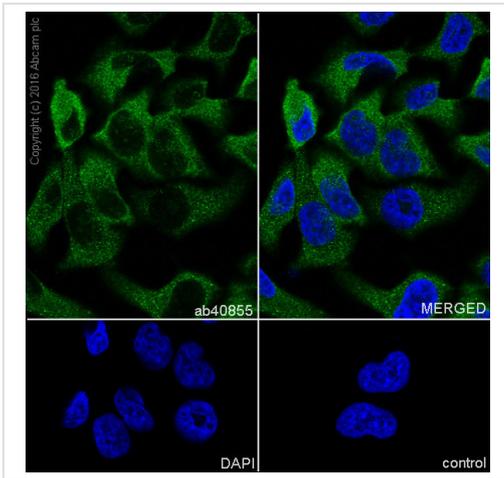


Flow Cytometry - Anti-Smad2 antibody [EP784Y]  
(ab40855)

ab40855 staining Smad2 in the human cell line HeLa (Human epithelial cell line from cervix adenocarcinoma) by flow cytometry. Cells were fixed with 4% paraformaldehyde and the sample was incubated with the primary antibody at a dilution of 1/20. A goat anti rabbit IgG (Alexa Fluor® 488) at a dilution of 1/2000 was used as the secondary antibody.

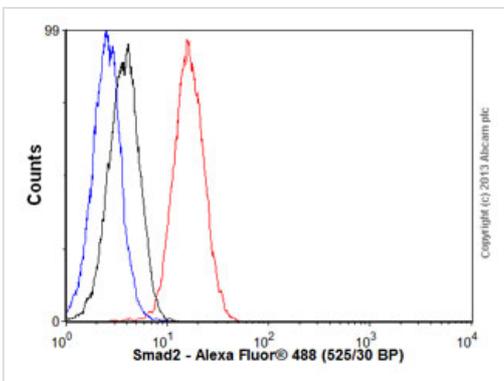
Isotype control: Rabbit monoclonal IgG (Black)

Unlabelled control: Cell without incubation with primary antibody and secondary antibody (Blue)



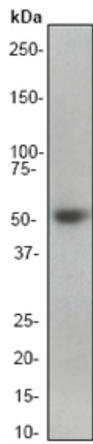
Immunocytochemistry/ Immunofluorescence - Anti-Smad2 antibody [EP784Y] (ab40855)

Immunofluorescence staining of HeLa cells with purified ab40855 at a working dilution of 1/500, counter-stained with DAPI. The secondary antibody was an Alexa Fluor® 488 conjugated goat anti-rabbit (ab150077), used at a dilution of 1/1000. The cells were fixed in 4% PFA and permeabilized using 0.1% Triton X 100. The negative control is shown in bottom right hand panel - for the negative control, PBS was used instead of the primary antibody.



Flow Cytometry - Anti-Smad2 antibody [EP784Y] (ab40855)

Overlay histogram showing PC3 cells stained with ab40855 (red line). The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab40855, 1/100 dilution) for 30 min at 22°C. The secondary antibody used was Alexa Fluor® 488 goat anti-rabbit IgG (H&L) (ab150077) at 1/2000 dilution for 30 min at 22°C. Isotype control antibody (black line) was rabbit IgG (monoclonal) (1µg/1x10<sup>6</sup> cells) used under the same conditions. Unlabelled sample (blue line) was also used as a control. Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 525/30 bandpass filter.

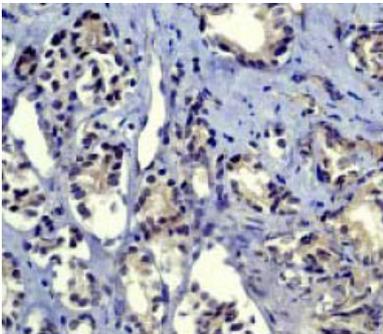


Anti-Smad2 antibody [EP784Y] (ab40855) at  
1/500000 dilution + Jurkat cell lysate

**Predicted band size** : 58 kDa

**Observed band size** : 58 kDa

Western blot - Smad2 antibody [EP784Y] (ab40855)



ab40855 at a 1:100 dilution staining Smad2 in  
human prostate carcinoma tissue.

Immunohistochemistry (Formalin/PFA-fixed paraffin-  
embedded sections) - Anti-Smad2 antibody  
[EP784Y] (ab40855)

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