# Phospho-SMAD1-S463/S465+SMAD5-S463/S465+SMAD9-S465/S467 Rabbit mAb

Catalog No.: AP1518 Recombinant

# **Basic Information**

Observed MW 60kDa

Calculated MW 52kDa

**Category** SMab Recombinant Monoclonal Antibody

Applications WB,IHC-P,IF/ICC,ELISA

**Cross-Reactivity** Human,Mouse,Rat

CloneNo number ARC59007

## **Recommended Dilutions**

WB	1:1000 - 1:2000
IHC-P	1:50 - 1:200
IF/ICC	1:100 - 1:400
ELISA	Recommended starting concentration is 1 µg/mL. Please optimize the concentration based on your specific assay requirements.

## Contact

www.abclonal.com

# Background

The protein encoded by this gene is involved in the transforming growth factor beta signaling pathway that results in an inhibition of the proliferation of hematopoietic progenitor cells. The encoded protein is activated by bone morphogenetic proteins type 1 receptor kinase, and may be involved in cancer. Alternative splicing results in multiple transcript variants.

# **Immunogen Information**

**Gene ID** 4086/4090/4093 Swiss Prot Q15797/Q99717/O15198

#### Immunogen

A synthetic phosphorylated peptide around S463/S465//S467 of human SMAD1/SMAD5/SMAD9.

## Synonyms

BSP1; MADH1; MADR1; JV4-1; DWFC; JV5-1; MADH5; MADH6; MADH9; SMAD8; Phospho-SMAD1-S463/S465+SMAD5-S463/S465+SMAD9-S465/S467

## **Product Information**

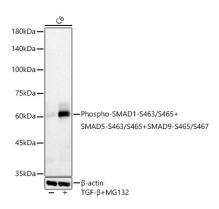
**Source** Rabbit **Isotype** IgG **Purification** Affinity purification

## Storage

Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.09% Sodium azide,0.05% BSA,50% glycerol,pH7.3.

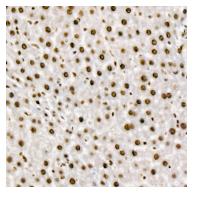


## Validation Data



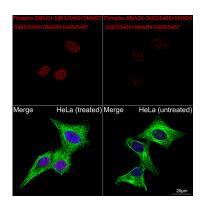
Western blot analysis of lysates from C6 cells using Phospho-SMAD1-S463/S465+SMAD5-S463/S465+SMAD9-S465/S467 Rabbit mAb (AP1518) at 1:1000 dilution. C6 cells were treated by TGF beta 3(10ng/ml) and MG132(2 µM) at 37°C for 20 minutes. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 25µg per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020).

Immunohistochemistry analysis of paraffin-embedded Human lung cancer tissue using Phospho-SMAD1-S463/S465+SMAD5-S463/S465+SMAD9-S465/S467 Rabbit mAb (AP1518) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate Bufferr (pH 6.0) prior to IHC staining.



Exposure time: 30s.

Immunohistochemistry analysis of paraffin-embedded Rat liver tissue using Phospho-SMAD1-S463/S465+SMAD5-S463/S465+SMAD9-S465/S467 Rabbit mAb (AP1518) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate Bufferr (pH 6.0) prior to IHC staining.



Confocal imaging of HeLa cells (treated with BMP2) and HeLa cells (untreated) using Phospho-SMAD1-S463/S465+SMAD5-\$463/\$465+\$MAD9-\$465/\$467 Rabbit mAb (AP1518, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). The cells were counterstained with  $\alpha$ tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.