



# SMAD2 Recombinant Rabbit Monoclonal Antibody (31H15L4)

<b>Product Details</b>	
Size	100 μg
Species Reactivity	Human
Published Species	Human
Host/Isotype	Rabbit / IgG
Expression system	Expi293
Class	Recombinant Monoclonal
Туре	Antibody
Clone	31H15L4
Conjugate	Unconjugated
Immunogen	A peptide corresponding to amino acids 81-107 of Q15796.
Form	Liquid
Concentration	0.5 mg/mL
Purification	Protein A
Storage buffer	PBS
Contains	0.09% sodium azide
Storage conditions	Store at 4°C short term. For long term storage, store at -20°C, avoiding freeze/thaw cycles.
RRID	AB_2532277

Applications	Tested Dilution	Publications
Western Blot (WB)	0.5-1 μg/mL	3 Publications
Immunohistochemistry (Paraffin) (IHC (P))	-	1 Publication
Immunocytochemistry (ICC/IF)	1 μg/mL	-
Flow Cytometry (Flow)	1-3 µg/1x10^6 cells	-
ELISA (ELISA)	1-5 μg/mL	-
ChIP assay (ChIP)	3 µg	1 Publication
Miscellaneous PubMed (Misc)	-	1 Publication

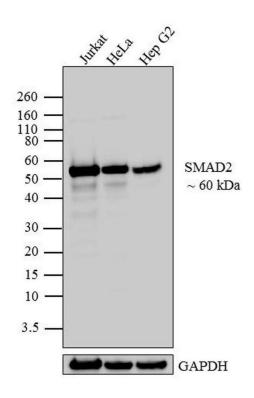
#### **Product Specific Information**

This antibody is predicted to react with bovine, carp, chicken, goldfish, mouse, opossum, orangutan, rat, Xenopus and zebrafish based on sequence homology.

Intact IgG appears on a non-reducing gel as ~150 kDa band and upon reduction generating a ~25 kDa light chain band and a ~50 kDa heavy chain.

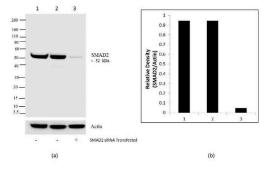
Recombinant rabbit monoclonal antibodies are produced using in vitro expression systems. The expression systems are developed by cloning in the specific antibody DNA sequences from immunoreactive rabbits. Then, individual clones are screened to select the best candidates for production. The advantages of using recombinant rabbit monoclonal antibodies include: better specificity and sensitivity, lot-to-lot consistency, animal origin-free formulations, and broader immunoreactivity to diverse targets due to larger rabbit immune repertoire.

## Product Images For SMAD2 Recombinant Rabbit Monoclonal Antibody (31H15L4)



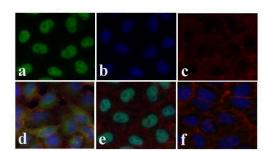
#### SMAD2 Antibody (700048) in WB

Western blot analysis of SMAD2 was performed by loading 20 µg of Jurkat (lane1), HeLa (lane2) and Hep G2 (lane3), cell lysates using Novex® NuPAGE® 4-12 % Bis-Tris gel (Product # NP0321BOX), XCell SureLock Electrophoresis System (Product # El0002), Novex® Sharp Pre-Stained Protein Standard (Product # LC5800), and iBlot® Dry Blotting System (Product # IB21001). Proteins were transferred to a nitrocellulose membrane and blocked with 5 % skim milk for 1 hour at room temperature. SMAD2 was detected at ~60 kDa using SMAD2 Recombinant Rabbit Monoclonal Antibody (Product # 700048) at 0.5-1 µg /mL in 2.5 % skim milk at 4°C overnight on a rocking platform. Goat anti-Rabbit IgG-HRP Secondary Antibody (Product # G-21234) at 1:5000 dilution was used and chemiluminescent detection was performed using Pierce™ ECL Western blotting Substrate (Product # 32106).



### SMAD2 Antibody (700048)

Antibody specificity was demonstrated by siRNA mediated knockdown of the target protein. HeLa cells were transfected with SMAD2 siRNA and decrease in signal intensity was observed in Western blot application (Fig a) using Anti-SMAD2 Recombinant Rabbit Monoclonal Antibody (31H15L4) (Product # 700048). Densitometric analysis of this Western blot is shown in histogram (Fig b). {KD}



#### SMAD2 Antibody (700048) in ICC/IF

Immunofluorescence analysis of SMAD2 was done on 70% confluent log phase TGF- beta treated HeLa cells (serum starved for 16 hours followed by treatment with 20 ng/mL TGF-beta for 1 hour). The cells were fixed with 4% paraformaldehyde for 15 minutes, permeabilized with 0.25% Triton X-100 for 10 minutes, and blocked with 5% BSA for 1 hour at room temperature. The cells were labeled with SMAD2 Recombinant Rabbit Monoclonal Antibody (Product # 700048) at 1 μg/mL in 1% BSA and incubated for 3 hours at room temperature and then labeled with Alexa Fluor 488 Goat anti-Rabbit IgG Secondary Antibody (Product # A-11008) at a dilution of 1:400 for 30 minutes at room temperature (Panel a: green). Nuclei (Panel b: blue) were stained with SlowFade Gold Antifade Mountant DAPI (Product # S36938). F-actin (Panel c: red) was stained with Alexa Fluor 594 Phalloidin (Product # A12381). Panel d is an untreated HeLa cells showing cytoplasmic localization. Panel e shows nuclear localization of SMAD2 upon treatment. Panel f is no primary antibody control. The images were captured at 20X magnification.

## View more figures on thermofisher.com

#### □ 6 References

## Western Blot (3)

Cancers

Sulforaphane Inhibits the Expression of Long Noncoding RNA H19 and Its Target APOBEC3G and Thereby Pancreatic Cancer Progression.

"Published figure using SMAD2 recombinant monoclonal antibody (Product # 700048) in Western Blot" Authors: Luo Y,Yan B,Liu L,Yin L,Ji H,An X,Gladkich J,Qi Z,De La Torre C,Herr I

**Year** 2021

Species Human

International journal of molecular medicine

Evaluation of the inhibitory effect of tacrolimus combined with mycophenolate mofetil on mesangial cell proliferation based on the cell cycle.

"Published figure using SMAD2 recombinant monoclonal antibody (Product # 700048) in Western Blot" Authors: Gao Y,Yang H,Wang Y,Tian J,Li R,Zhou X

**Year** 2020

Species Human

Dilution 1:200

View more WB references on thermofisher.com

# Immunohistochemistry (Paraffin) (1)

Molecular endocrinology (Baltimore, Md.)

FGFR2IIIb-MAPK Activity Is Required for Epithelial Cell Fate Decision in the Lower Müllerian Duct.

"Published figure using SMAD2 recombinant monoclonal antibody (Product # 700048) in Immunofluorescence" Authors: Terakawa J,Rocchi A,Serna VA,Bottinger EP,Graff JM,Kurita T

**Year** 2016

Dilution 1:100

## ChIP assay (1)

Developmental cell

Long-Range Signaling Activation and Local Inhibition Separate the Mesoderm and Endoderm Lineages.

"Published figure using SMAD2 monoclonal antibody (Product # 700048) in ChIP assay" Authors: van Boxtel AL.Economou AD.Heliot C.Hill CS

**Year** 2018

Species Human

# More applications with references on thermofisher.com

## Misc (1)

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