

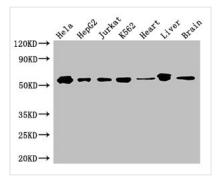
Image





SMAD2 Antibody

Product Code	CSB-PA15557A0Rb
Storage	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
Uniprot No.	Q15796
Immunogen	Recombinant Human Mothers against decapentaplegic homolog 2 protein (2-467AA)
Raised In	Rabbit
Species Reactivity	Human, Mouse
Tested Applications	ELISA, WB, IHC, IF; Recommended dilution: WB:1:500-1:5000, IHC:1:100-1:1000, IF:1:50-1:500
Form	Liquid
Conjugate	Non-conjugated
Storage Buffer	Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, PH 7.4
Purification Method	>95%, Protein G purified
Isotype	IgG
Clonality	Polyclonal
Alias	Mothers against decapentaplegic homolog 2 (MAD homolog 2) (Mothers against DPP homolog 2) (JV18-1) (Mad-related protein 2) (hMAD-2) (SMAD family member 2) (SMAD 2) (Smad2) (hSMAD2), SMAD2, MADH2 MADR2
Immunogen Species	Homo sapiens (Human)
Research Area	Signal Transduction
Target Names	SMAD2



Western Blot

Positive WB detected in: Hela whole cell lysate, HepG2 whole cell lysate, Jurkat whole cell lysate, K562 whole cell lysate, Mouse heart tissue, Mouse liver tissue, Mouse brain tissue All lanes: SMAD2 antibody at 3μg/ml

Secondary

Goat polyclonal to rabbit IgG at 1/50000 dilution

Predicted band size: 53, 49 kDa Observed band size: 53 kDa

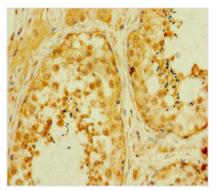




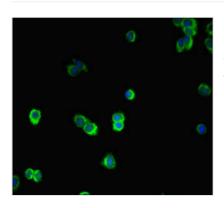




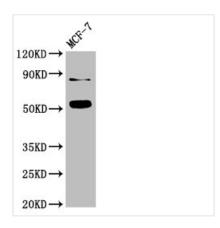




Immunohistochemistry of paraffin-embedded human testis tissue using CSB-PA15557A0Rb at dilution of 1:100



Immunofluorescent analysis of HepG2 cells using CSB-PA15557A0Rb at dilution of 1:100 and Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L)



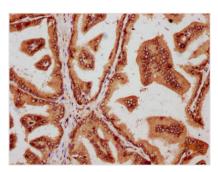
Western Blot

Positive WB detected in: MCF-7 whole cell lysate

All lanes: SMAD2 antibody at 3µg/ml

Goat polyclonal to rabbit IgG at 1/50000 dilution

Predicted band size: 53, 49 kDa Observed band size: 53 kDa



IHC image of CSB-PA15557A0Rb diluted at 1:600 and staining in paraffin-embedded human prostate tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.

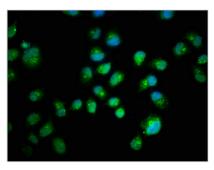


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Immunofluorescence staining of Hela cells with CSB-PA15557A0Rb at 1:200, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).