

BMP1 Antibody

Cat.#: DF9280 Concn.: 1mg/ml Mol.Wt.: 111 kDa, 95 kDa Size: 100ul,200ul Source: Rabbit Clonality: Polyclonal

Application: WB 1:1000-3000, IHC 1:50-1:200, ELISA(peptide)

1:20000-1:40000

*The optimal dilutions should be determined by the end

user.

Reactivity: Human, Mouse, Rat

Purification: The antiserum was purified by peptide affinity

chromatography using SulfoLink™ Coupling Resin (Thermo

Fisher Scientific).

Specificity: BMP1 Antibody detects endogenous levels of total BMP1.

Immunogen: A synthesized peptide derived from human BMP1,

corresponding to a region within the internal amino acids.

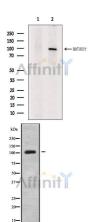
Uniprot: P13497

Storage Condition and

Buffer:

Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at -20

°C.Stable for 12 months from date of receipt.



Western blot analysis of extracts from C6, using BMP1 Antibody. The lane on the left was treated with blocking peptide.

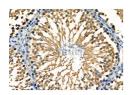
Observed bands: 95 kDa.

Western blot analysis of BMP1 using Jurkat whole cell lysates

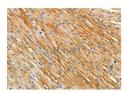


Affinity Biosciences website:www.affbiotech.com

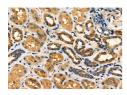
website:www.affbiotech.com order:order@affbiotech.com



DF9280 at 1/100 staining Rat testis tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the primary antibody at 4° C overnight. An HRP conjugated anti-Rabbit antibody was used as the secondary antibody.



DF9280 at 1/100 staining Rat heart tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the primary antibody at 4°C overnight. An HRP conjugated anti-Rabbit antibody was used as the secondary antibody.



DF9280 at 1/100 staining Human kidney cancer and adjacent normal tissues by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the primary antibody at 4°C overnight. An HRP conjugated anti-Rabbit antibody was used as the secondary antibody.

<code>IMPORTANT:</code> For western blot, incubate membrane with diluted primary Ab in 5% w/v milk , 1X TBS, 0.1% Tween\$20 at 4°C with gentle shaking, overnight.

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