

## Catalog No. BN41631R

## Rabbit Anti-CD86 Polyclonal Antibody

DATASHEET	
Host:Rabbit	Concentration:1mg/ml
Target Protein:CD86	Applications:WB(1:500-2000) ELISA(1:5000-10000)
IR:Immunogen Range:140-175/313	IHC-P(1:100-500)
Clonality:Polyclonal	IF(1:100-500)
Isotype:IgG	Cross Reactive Species:Human
Entrez Gene: 56822	Rat
Swiss Prot: <u>O35531</u>	Dog Pig
<b>Source:</b> KLH conjugated synthetic peptide derived from the middle of rat CD86:140-175/313	Cow Sheep
Purification: affinity purified by Protein A	For research use only. Not intended for
<b>Storage:</b> 0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol. Shipped at $4^{\circ}$ C. Store at -20 °C for one year. Avoid repeated freeze/thaw cycles.	diagnostic or therapeutic use.
<b>Background:</b> This gene encodes a type I membrane protein that is a member of the immunoglobulin superfamily. This protein is expressed by antigen-presenting cells, and it is the ligand for two proteins at the cell surface of T cells, CD28 antigen and cytotoxic T-lymphocyte-associated protein 4. Binding of this protein with CD28 antigen is a costimulatory signal for activation of the T-cell. Binding of this protein with cytotoxic T-lymphocyte-associated protein 4 negatively regulates T-cell activation and diminishes the immune response. Alternative splicing results in several transcript variants encoding different isoforms.[provided by RefSeq, May 2011].	



## VALIDATION IMAGES



Sample:

Lane 1: HepG2 (Human) Cell Lysate at 30 ug Lane 2: U937 (Human) Cell Lysate at 30 ug Lane 3: Spleen (Rat) Lysate at 40 ug Lane 4: Spleen (Mouse) Lysate at 40 ug Lane 5: HL-60 (Human) Cell Lysate at 30 ug Primary: Anti-CD86 at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 72-74 kD Observed band size: 72 kD

Sample:

Raji(Human) Cell Lysate at 30 ug HepG2(Human) Cell Lysate at 30 ug Primary: Anti- CD86 at 1/300 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 70/80 kD Observed band size: 70 kD



Tissue/cell: rat lung tissue; 4% Paraformaldehyde-fixed and paraffin-embedded;

Antigen retrieval: citrate buffer ( 0.01M, pH 6.0 ), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum) at 37°C for 20 min;

Incubation: Anti-CD86/B7-2 Polyclonal Antibody, Unconjugated 1:200, overnight at 4°C, followed by conjugation to the secondary antibody and DAB staining







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Tissue/cell: Human esophageal carcinoma; 4% Paraformaldehyde-fixed and paraffin-embedded; Antigen retrieval: citrate buffer ( 0.01M, pH 6.0 ), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum) at 37 °C for 20 min;

Incubation: Anti-CD86/B7-2 Polyclonal Antibody, Unconjugated 1:200, overnight at 4°C, followed by conjugation to the secondary antibody and DAB staining

Blank control: Mouse spleen.

Primary Antibody (green line): Rabbit Anti-CD86 antibody Dilution: 2µg /10^6 cells;

Isotype Control Antibody (orange line): Rabbit IgG . Protocol

The cells were incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at at room temperature .Cells stained with Primary Antibody for 30 min at room temperature. Acquisition of 20,000 events was performed.

Blank control: U937(blue).

Primary Antibody: Rabbit Anti-CD86 antibody, Dilution: 1µg in 100 µL 1X PBS containing 0.5% BSA;

Isotype Control Antibody: Rabbit IgG (orange) ,used under the same conditions.

Secondary Antibody: Goat anti-rabbit IgG-PE(white blue), Dilution: 1:200 in 1 X PBS containing 0.5% BSA.

Protocol

The cells were fixed with 2% paraformaldehyde (10 min).Primary antibody (1 $\mu$ g /1x10^6 cells) were incubated for 30 min on the ice, followed by 1 X PBS containing 0.5% BSA + 10% goat serum (15 min) to block non-specific protein-protein interactions. Then the Goat Anti-rabbit IgG/PE antibody was added into the blocking buffer mentioned above to react with the primary antibody at 1/200 dilution for 30 min on ice. Acquisition of 20,000 events was performed.