

## Anti-Bad Rabbit mAb

Purified Recombinant Rabbit Monoclonal Antibody

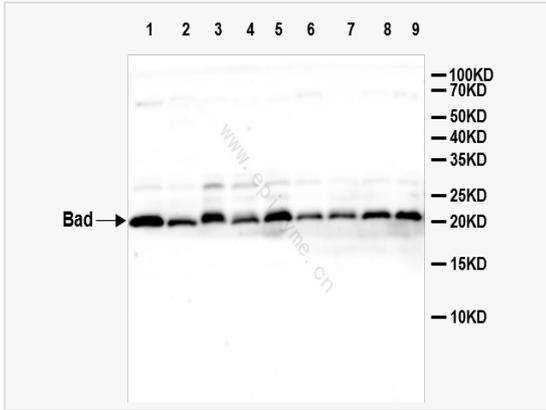
Catalog # R013754

### Product Information

|             |   |
|-------------|---|
| Application | ELISA, IHC-P/IF (Tissue-P), WB, IF (Cell)/ICC   |
| Reactivity  | Mouse, Rat, Human   |
| Dilution    | WB 1:1,000~1:2,000; IHC-P 1:200; IF 1:100   |
| Host        | Rabbit  |
| Clonality   | Monoclonal  |
| Clone No.   | 39D84M92  |
| Isotype     | IgG   |
| Label       | Unconjugated  |
| Immunogen   | A synthesized peptide derived from human Bad  |
| Format      | Affinity purified monoclonal antibody supplied in PBS with 0.01% sodium azide and 50% glycerol, pH 7.3.                     |
| Storage     | Shipped on wet ice. Store at -20°C. Stable for 24 months from date of receipt. Aliquoting is unnecessary for -20°C storage. |
| Precautions | Anti-Bad Rabbit mAb [39D84M92] is for research use only and not for use in diagnostic or therapeutic procedures.            |

### Protein Information

|                   |   |
|-------------------|---|
| Synonyms          | BAD, BBC6, BCL2L8, Bcl2 antagonist of cell death, BAD, Bcl-2-binding component 6, Bcl-2-like protein 8, Bcl2-L-8, Bcl-XL/Bcl-2-associated death promoter.   |
| Calculated MW     | Calculated MW: 18 kDa; Observed MW: 18 kDa  |
| Uniprot ID        | Q92934  |
| Gene ID           | 572   |
| Background        | Promotes cell death. Successfully competes for the binding to Bcl-X(L), Bcl-2 and Bcl-W, thereby affecting the level of heterodimerization of these proteins with BAX. Can reverse the death repressor activity of Bcl-X(L), but not that of Bcl-2 (By similarity). Appears to act as a link between growth factor receptor signaling and the apoptotic pathways. |
| Cellular Location | Mitochondrion outer membrane. Cytoplasm. Note=Colocalizes with HIF3A isoform 2 in the cytoplasm (PubMed:21546903). Upon phosphorylation, locates to the cytoplasm.  |



Western Blot - Anti-Bad Rabbit mAb [39D84M92]

All lanes: R013754 at 1:1,000 dilution

Lane 1: HeLa (human cervix adenocarcinoma epithelial cell) whole cell lysates

Lane 2: MCF7 (human breast adenocarcinoma epithelial cell) whole cell lysates

Lane 3: Jurkat (human T lymphocytic leukemia cell) whole cell lysates

Lane 4: HCT116 (human colorectal carcinoma epithelial cell) whole cell lysates

Lane 5: T24 (human bladder cancer epithelial cell) whole cell lysates

Lane 6: HepG2 (human hepatocellular carcinoma epithelial cell) whole cell lysates

Lane 7: SCC-9 (human tongue squamous carcinoma epithelial cell) whole cell lysates

Lane 8: U2OS (human osteosarcoma epithelial cell) whole cell lysates

Lane 9: SW620 (human colorectal carcinoma epithelial cell) whole cell lysates

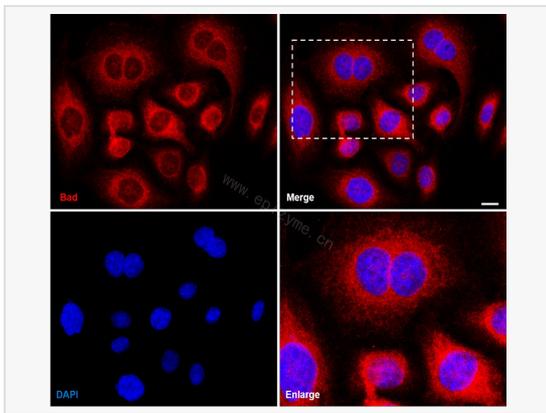
Lysates/proteins at 10 µg per lane.

Secondary antibody: Goat Anti-Rabbit IgG(H+L), HRP Conjugated (Cat. No. LF102) at 1:5,000 dilution

Predicted band size: 18 kDa

Observed band size: 18 kDa

Developed using the ECL technique (Cat. No. SQ201).



Immunofluorescence - Anti-Bad Rabbit mAb [39D84M92]

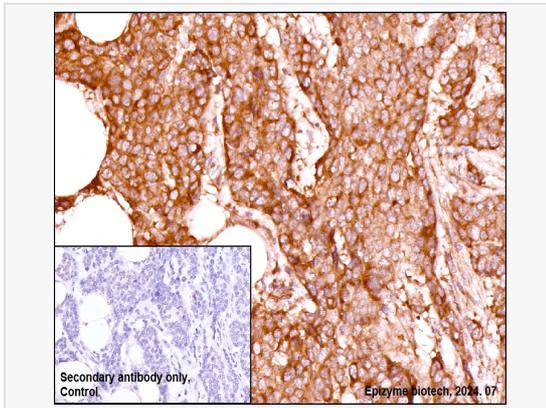
Sample: HeLa cells

The cells were fixed with 4% paraformaldehyde (10 min), permeabilized with 0.5% Triton X-100 for 10 minutes and then blocked with 5% BSA in 0.1% PBS-Tween for 0.5 hours.

Primary antibody: R013754 at 1:100 dilution

Secondary antibody: Goat anti-Rabbit (555) at 1:1,000 dilution (shown in red)

Nuclei were stained with DAPI (shown in blue).



Immunohistochemistry - Anti-Bad Rabbit mAb [39D84M92]

Sample: Paraformaldehyde-fixed, paraffin embedded human breast carcinoma tissue

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0) for 30 mins.

Primary antibody: R013754 at 1:200 dilution

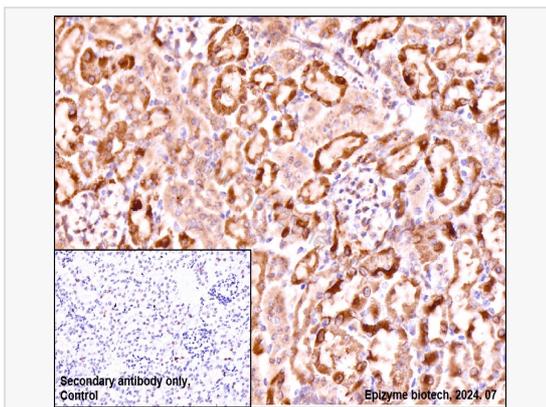
Secondary antibody: Goat Anti-Rabbit IgG (H+L), HRP conjugated at 1:1,000 dilution

DAB was used as the chromogen.

Counter stained with hematoxylin.

Positive/negative staining were presented.

Only the secondary antibody was used as the negative control.



Immunohistochemistry - Anti-Bad Rabbit mAb [39D84M92]

Sample: Paraformaldehyde-fixed, paraffin embedded mouse kidney tissue

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0) for 30 mins.

Primary antibody: R013754 at 1:200 dilution

Secondary antibody: Goat Anti-Rabbit IgG (H+L), HRP conjugated at 1:1,000 dilution

DAB was used as the chromogen.

Counter stained with hematoxylin.

Positive/negative staining were presented.

Only the secondary antibody was used as the negative control.