

A21902

Leader in Biomolecular Solutions for Life Science



## [KO Validated] DDIT3/CHOP Rabbit mAb

Catalog No.: A21902

**KO** Validated

Recombinant

4 Publications

### Basic Information

#### Observed MW

27 kDa

#### Calculated MW

19 kDa

#### Category

SMab Recombinant Monoclonal  
Antibody

#### Applications

WB,IHC-P,IF/ICC,ELISA

#### Cross-Reactivity

Human,Mouse,Rat

#### CloneNo number

ARC51417

### Background

This gene encodes a member of the CCAAT/enhancer-binding protein (C/EBP) family of transcription factors. The protein functions as a dominant-negative inhibitor by forming heterodimers with other C/EBP members, such as C/EBP and LAP (liver activator protein), and preventing their DNA binding activity. The protein is implicated in adipogenesis and erythropoiesis, is activated by endoplasmic reticulum stress, and promotes apoptosis. Fusion of this gene and FUS on chromosome 16 or EWSR1 on chromosome 22 induced by translocation generates chimeric proteins in myxoid liposarcomas or Ewing sarcoma. Multiple alternatively spliced transcript variants encoding two isoforms with different length have been identified.

### Recommended Dilutions

**WB** 1:1000 - 1:2000

**IHC-P** 1:100 - 1:1000

**IF/ICC** 1:50 - 1:200

**ELISA** Recommended starting concentration is 1 µg/mL. Please optimize the concentration based on your specific assay requirements.

### Immunogen Information

#### Gene ID

1649

#### Swiss Prot

P35638

#### Immunogen

Synthetic peptide. This information is considered to be commercially sensitive.

#### Synonyms

CHOP; CEBPZ; CHOP10; CHOP-10; GADD153; AltDDIT3; C/EBPzeta; DDIT3/CHOP

### Product Information

#### Source

Rabbit

#### Isotype

IgG

#### Purification

Affinity purification

#### Storage

Store at -20°C. Avoid freeze / thaw cycles.

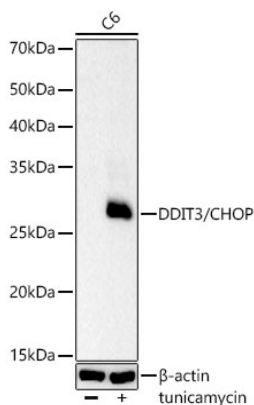
Buffer: PBS containing 50% glycerol and 0.05% BSA, preserved with proclin300 or sodium azide (as specified on the Certificate of Analysis), pH 7.3.

### Contact



[www.abclonal.com](http://www.abclonal.com)

## Validation Data



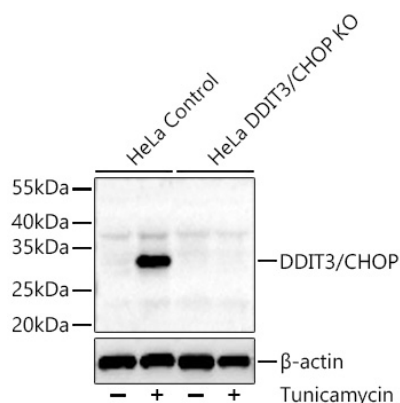
Western blot analysis of lysates from C6 cells using [KO Validated] DDIT3/CHOP Rabbit mAb (A21902) at 1:900 dilution incubated overnight at 4°C. C6 cells were treated with tunicamycin (2 µg/ml) for 8 hours  
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25 µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 60s.



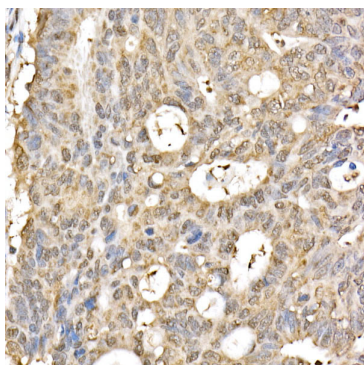
Western blot analysis of lysates from wild type (WT) and DDIT3/CHOP knockout (KO) HeLa cells using DDIT3/CHOP Rabbit mAb (A21902) at 1:1000 dilution incubated overnight at 4°C. HeLa cells were treated with Tunicamycin (20 µg/mL) at 37°C for 3 hrs. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25 µg per lane.

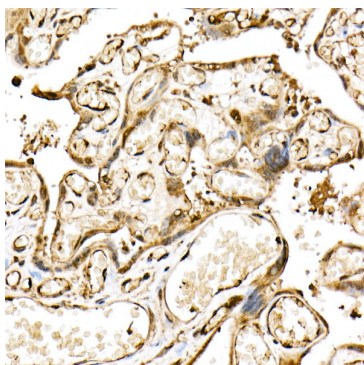
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

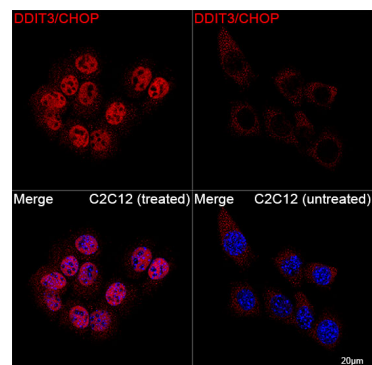
Exposure time: 90s.



Immunohistochemistry analysis of paraffin-embedded Human colon carcinoma tissue using [KO Validated] DDIT3/CHOP Rabbit mAb (A21902) at a dilution of 1:100 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate Buffer (pH 6.0) prior to IHC staining.

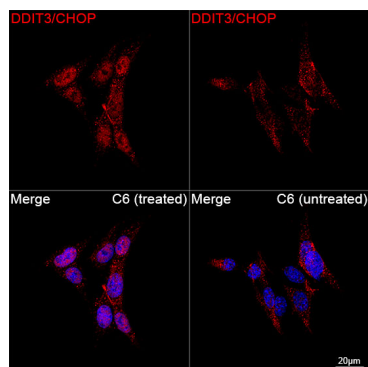


Immunohistochemistry analysis of paraffin-embedded Human placenta tissue using [KO Validated] DDIT3/CHOP Rabbit mAb (A21902) at a dilution of 1:100 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate Buffer (pH 6.0) prior to IHC staining.

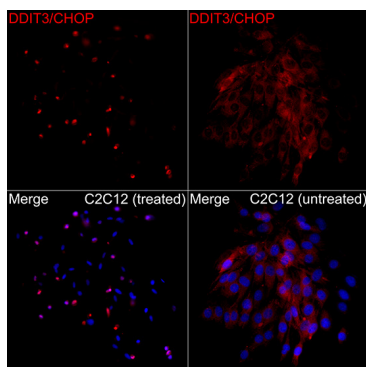


Confocal imaging of C2C12 cells (treated with Tunicamycin) and C2C12 cells (untreated) cells using [KO Validated] DDIT3/CHOP Rabbit mAb(A21902, dilution 1:200) followed by a further incubation with Cy3-conjugated Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). DAPI was used for nuclear staining (Blue). Objective: 100x.

## Validation Data



Confocal imaging of C6 cells (treated with Tunicamycin) and C6 cells (untreated) using [KO Validated] DDIT3/CHOP Rabbit mAb (A21902, dilution 1:200) followed by a further incubation with Cy3-conjugated Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). DAPI was used for nuclear staining (Blue). Objective: 100x.



Immunofluorescence analysis of C2C12 cells (treated with Tunicamycin) and C2C12 cells (untreated) using [KO Validated] DDIT3/CHOP Rabbit mAb (A21902) at a dilution of 1:200 (40x lens). Secondary antibody: Cy3 Goat anti-Rabbit IgG (H+L) (AS007) at 1:500 dilution. Blue: DAPI for nuclear staining.