

## CD40 (Phospho Thr254) Rabbit pAb

CatalogNo: YP0379 **Orthogonal Validated** 

### Key Features

#### Host Species

- Rabbit

#### Reactivity

- Human, Mouse

#### Applications

- WB, ELISA

#### MW

- 30kD (Observed)

#### Isotype

- IgG

### Storage

**Storage\*** -15°C to -25°C/1 year (Do not lower than -25°C)

**Formulation** Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

### Recommended Dilution Ratios

**WB 1:500-1:2000**

**ELISA 1:5000**

**Not yet tested in other applications.**

### Basic Information

**Clonality** Polyclonal

### Immunogen Information

**Immunogen** The antiserum was produced against synthesized peptide derived from human TNFRSF5 around the phosphorylation site of Thr254. AA range:220-269

**Specificity** Phospho-CD40 (T254) Polyclonal Antibody detects endogenous levels of CD40 protein only when phosphorylated at T254. The name of modified sites may be influenced by many factors, such as species (the modified site was not originally found in human samples) and the change of protein sequence (the previous protein sequence is incomplete, and the protein sequence may be prolonged with the development of protein sequencing technology). When naming, we will use the "numbers" in historical reference to keep the sites consistent with the reports. The antibody binds to the following modification sequence (lowercase letters are modification sites): QETLH

## Target Information

**Gene name** CD40

**Protein Name** Tumor necrosis factor receptor superfamily member 5

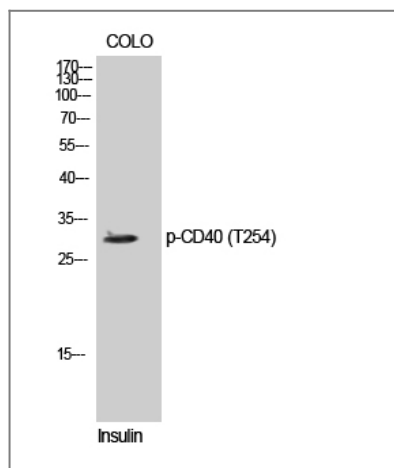
Organism	Gene ID	UniProt ID
Human	<a href="#">958</a> ;	<a href="#">P25942</a> ;
Mouse	<a href="#">21939</a> ;	<a href="#">P27512</a> ;

**Cellular Localization** [Isoform I]: Cell membrane; Single-pass type I membrane protein.; [Isoform II]: Secreted.

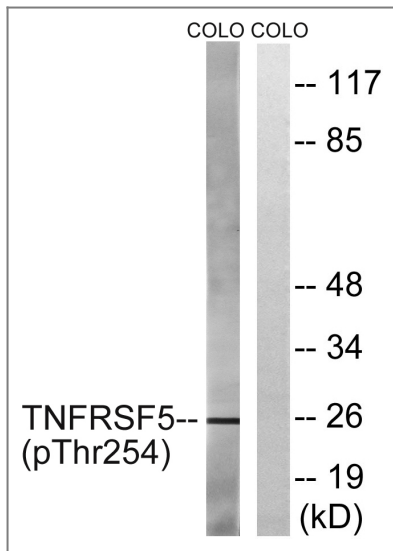
**Tissue specificity** B-cells and in primary carcinomas.

**Function** Alternative products:Additional isoforms seem to exist,Disease:Defects in CD40 are the cause of hyper-IgM immunodeficiency type 3 (HIGM3) [MIM:606843]. HIGM3 is an autosomal recessive disorder which includes an inability of B cells to undergo isotype switching, one of the final differentiation steps in the humoral immune system, an inability to mount an antibody-specific immune response, and a lack of germinal center formation.,Function:Receptor for TNFSF5/CD40LG.,online information:CD40 entry,online information:CD40 mutation db,similarity:Contains 4 TNFR-Cys repeats.,subunit:Monomer and homodimer. The variant form found in the bladder carcinoma cell line Hu549 does not form homodimers. Interacts with TRAF1, TRAF2, TRAF3, TRAF5 and TRAF6.,tissue specificity:B-cells and in primary carcinomas.,

## Validation Data



Western Blot analysis of COLO cells using Phospho-CD40 (T254) Polyclonal Antibody diluted at 1:500



Western blot analysis of lysates from COLO205 cells treated with Insulin 0.01U/ml 15', using TNFRSF5 (Phospho-Thr254) Antibody. The lane on the right is blocked with the phospho peptide.

## Contact information

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**CD40 (Phospho Thr254) Rabbit pAb**

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