

## Caspase-1 (AB2378) mouse mAb

W0179

### Key Features

#### Host Species

- Mouse

#### Reactivity

- Human; Mouse; Rat; Horse; Pig;

#### Applications

- WB; IF; ELISA

#### MW

- 45 kDa (calculated)
- 45 kDa (observed)

#### Isotype

- IgG2b, kappa

### Recommended Dilution Ratios

#### Application

WB; IF; ELISA

#### Dilution

WB, 1:500-1:2000 | IF, 1:100-1:500 | ELISA, Recommended starting concentration is 1 µg/mL.

Please optimize the concentration based on your specific assay requirements.

### Storage

#### Storage Conditions

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

#### Storage buffer

The antibody is provided in liquid form in phosphate - buffered saline with 50% glycerol, 0.05% BSA, and 0.05% Proclin 300.

### Basic Information

**Clonality** Monoclonal

**Clone Number** AB2378

**Immunogen** A synthetic peptide corresponding to the amino acid region 350 - 404 of the human Caspase - 1 protein.

**Specificity** This antibody detects endogenous levels of Caspase-1 protein.

**Purification** Affinity purification Protein A

**Concentration** Product concentration may vary by batch. Please refer to the product COA for details.

### Target Information

**Gene name** CASP1

**Protein Name** Caspase-1

| Database Link | Organism | Swiss Prot. | Gene ID |
|---------------|----------|-------------|---------|
|               | Human    | P29466      | 834     |

#### Background

**caspase 1 (CASP1) Homo sapiens** This gene encodes a protein which is a member of the cysteine-aspartic acid protease (caspase) family. Sequential activation of caspases plays a central role in the execution-phase of cell apoptosis. Caspases exist as inactive proenzymes which undergo proteolytic processing at conserved aspartic residues to produce 2 subunits, large and small, that dimerize to form the active enzyme. This gene was identified by its ability to proteolytically cleave and activate the inactive precursor of interleukin-1, a cytokine involved in the processes such as inflammation, septic shock, and wound healing. This gene has been shown to induce cell apoptosis and may function in various developmental stages. Studies of a similar gene in mouse suggest a role in the pathogenesis of Huntington disease. Alternative splicing results in transcript variants encoding distinct isoforms.

W0179-V.2-EN

[provided by RefSeq, Mar 2012],