

HTRA2 Mouse mAb

CatalogNo: YM1524

| Key Features

Host Species

- Mouse

Reactivity

- Mouse, Hamster

Applications

- WB

MW

- 36kD (Observed)

| Recommended Dilution Ratios

WB 1:1000

| 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.

| Basic Information

Clonality Monoclonal

Clone Number 3G6

| Immunogen Information

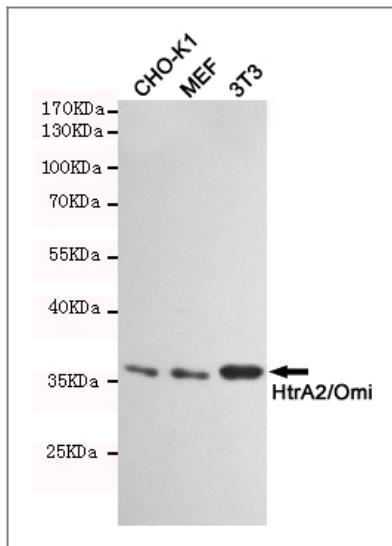
Immunogen Recombinant human HtrA2/Omi protein.

Specificity This antibody detects endogenous levels of HtrA2/Omi and does not cross-react with related proteins.

| Target Information

Gene name	HTRA2		
Protein Name	Serine protease HTRA2 mitochondrial		
	Organism	Gene ID	UniProt ID
	Human	27429 ;	O43464 ;
Cellular Localization	Mitochondrion intermembrane space. Mitochondrion membrane ; Single-pass membrane protein . Predominantly present in the intermembrane space. Released into the cytosol following apoptotic stimuli, such as UV treatment, and stimulation of mitochondria with caspase-8 truncated BID/tBID.; [Isoform 1]: Endoplasmic reticulum .		
Tissue specificity	[Isoform 1]: Ubiquitously expressed.		
Function	Catalytic activity: Cleavage of non-polar aliphatic amino-acids at the P1 position, with a preference for Val, Ile and Met. At the P2 and P3 positions, Arg is selected most strongly with a secondary preference for other hydrophilic residues., Disease: Defects in HTRA2 are the cause of Parkinson disease type 13 (PARK13) [MIM:610297, 168600]. Parkinson disease (PD) is a complex, multifactorial disorder that typically manifests after the age of 50 years, although early-onset cases (before 50 years) are known. PD generally arises as a sporadic condition but is occasionally inherited as a simple mendelian trait. Although sporadic and familial PD are very similar, inherited forms of the disease usually begin at earlier ages and are associated with atypical clinical features. PD is characterized by bradykinesia, resting tremor, muscular rigidity and postural instability, as well as by a clinically significant response to treatment with levodopa. The pathology involves the loss of dopaminergic neurons in the substantia nigra and the presence of Lewy bodies (intraneuronal accumulations of aggregated proteins), in surviving neurons in various areas of the brain., Domain: The mature N-terminus is involved in the interaction with XIAP., Domain: The PDZ domain mediates interaction with MXI2., Function: Serine protease that shows proteolytic activity against a non-specific substrate beta-casein. Promotes or induces cell death either by direct binding to and inhibition of BIRC proteins (also called inhibitor of apoptosis proteins, IAPs), leading to an increase in caspase activity, or by a BIRC inhibition-independent, caspase-independent and serine protease activity-dependent mechanism. Isoform 2 seems to be proteolytically inactive., PTM: Autoproteolytically activated., similarity: Belongs to the peptidase S1B family., similarity: Contains 1 PDZ (DHR) domain., subcellular location: Predominantly present in the intermembrane space. Released into the cytosol following apoptotic stimuli, such as UV treatment, and stimulation of mitochondria with caspase-8 truncated BID/tBID., subunit: Homotrimer. Interacts with MXI2. The mature protein, but not the precursor, binds to BIRC2, BIRC3 and XIAP., tissue specificity: Isoform 1 is ubiquitous; isoform 2 is expressed predominantly in the kidney, colon and thyroid.,		

| Validation Data



Western blot detection of HtrA2/Omi in CHO-K1,MEF and 3T3 cell lysates using HtrA2/Omi mouse mAb(dilution 1:1000).Predicted band size:49kDa.Observed band size:36kDa.

Contact information

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HTRA2 Mouse mAb

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