

# HSPD1 Antibody (C-term)

Purified Rabbit Polyclonal Antibody (Pab) Catalog # AW5439

## **Product Information**

Application	FC, IHC-P, WB
Primary Accession	<u>P10809</u>
Other Accession	<u>P63039, P63038, P18687, Q5ZL72, P31081</u>
Reactivity	Mouse, Rat, Human
Host	Rabbit
Clonality	Polyclonal
Calculated MW	61055
Isotype	Rabbit IgG
Antigen Source	HUMAN

## **Additional Information**

Gene ID	3329
Antigen Region	396-423
Other Names	60 kDa heat shock protein, mitochondrial, 60 kDa chaperonin, Chaperonin 60, CPN60, Heat shock protein 60, HSP-60, Hsp60, HuCHA60, Mitochondrial matrix protein P1, P60 lymphocyte protein, HSPD1, HSP60
Dilution	FC~~1:10~50 IHC-P~~1:100~500 WB~~1:1000
Target/Specificity	This HSPD1 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 396-423 amino acids from the C-terminal region of human HSPD1.
Format	Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is prepared by Saturated Ammonium Sulfate (SAS) precipitation followed by dialysis against PBS.
Storage	Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.
Precautions	HSPD1 Antibody (C-term) is for research use only and not for use in diagnostic or therapeutic procedures.

#### **Protein Information**

Name	HSPD1
Synonyms	HSP60

Function	Chaperonin implicated in mitochondrial protein import and macromolecular assembly. Together with Hsp10, facilitates the correct folding of imported proteins. May also prevent misfolding and promote the refolding and proper assembly of unfolded polypeptides generated under stress conditions in the mitochondrial matrix (PubMed: <u>11422376</u> , PubMed: <u>1346131</u> ). The functional units of these chaperonins consist of heptameric rings of the large subunit Hsp60, which function as a back- to-back double ring. In a cyclic reaction, Hsp60 ring complexes bind one unfolded substrate protein per ring, followed by the binding of ATP and association with 2 heptameric rings of the co-chaperonin Hsp10. This leads to sequestration of the substrate protein in the inner cavity of Hsp60 where, for a certain period of time, it can fold undisturbed by other cell components. Synchronous hydrolysis of ATP in all Hsp60 subunits results in the dissociation of the chaperonin rings and the release of ADP and the folded substrate protein (Probable).
Cellular Location	Mitochondrion matrix.

## Background

HSPD1 is a member of the chaperonin family. This protein may function as a signaling molecule in the innate immune system. The protein is essential for the folding and assembly of newly imported proteins in the mitochondria. The protein is adjacent to a related family member and the region between the 2 genes functions as a bidirectional promoter.

# References

Venner T.J., Singh B., Gupta R.S.DNA Cell Biol. 9:545-552(1990) Hansen J.J., Bross P., Westergaard M., Nielsen M.N., Eiberg H.,Hum. Genet. 112:71-77(2003) Rasmussen R.K., Ji H., Eddes J.S., Moritz R.L.,Electrophoresis 18:588-598(1997) Aboulaich N., Vainonen J.P., Stralfors P., Vener A.V.Biochem. J. 383:237-248(2004)

## Images



All lanes : Anti-HSPD1 Antibody (C-term) at 1:1000 dilution Lane 1: A431 whole cell lysates Lane 2: mouse liver lysates Lane 3: NIH/3T3 whole cell lysates Lane 4: rat liver lysates Lane 5: Hela whole cell lysates Lane 6: MOLT-4 whole cell lysates Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Rabbit IgG, (H+L),Peroxidase conjugated at 1/10000 dilution Predicted band size : 61 kDa Blocking/Dilution buffer: 5% NFDM/TBST.

Formalin-fixed and paraffin-embedded human lung carcinoma reacted with HSPD1 Antibody (C-term), which was peroxidase-conjugated to the secondary antibody, followed by DAB staining. This data demonstrates the use of this antibody for immunohistochemistry; clinical relevance has not been evaluated.





HSPD1 Antibody (C-term) (Cat. #AW5439) flow cytometric analysis of WiDr cells (right histogram) compared to a negative control cell (left histogram).FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis.

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