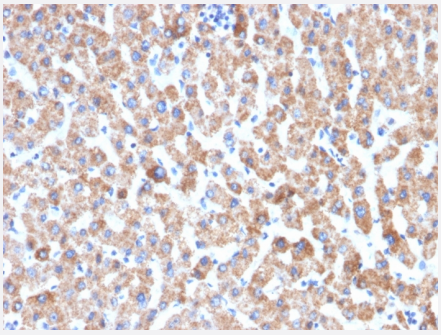
RecomAb™ Sm Size

# HSPD1 recombinant monoclonal antibody, clone HSPD1/2206R

Catalog # RAB08925 Size 20 ug

## Applications



### Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Immunohistochemical staining of human liver.




### Western Blot (Cell lysate)

Western Blot (cell lysate) analysis of HeLa cell lysate.

## Specification

| | |
|---|---|
| **Product Description** | Rabbit recombinant monoclonal antibody raised against full length human HSPD1. |
| **Antibody Species** | Rabbit |
| **Immunogen** | Recombinant protein corresponding to full-length human HSPD1. |
| **Reactivity** | Bovine, Chicken, Dog, Hamster, Human, Monkey, Mouse, Pig, Rabbit, Rat, Sheep |
| **Form** | Liquid |
| **Purification** | Protein A/G purification |

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| | |
|---|---|
| **Isotype** | IgG |
| **Recommend Usage** | Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) (0.5-1 ug/mL)<br>Western Blot (0.25-0.5 ug/mL)<br>The optimal working dilution should be determined by the end user. |
| **Storage Buffer** | In 10 mM PBS (0.05% BSA and 0.05% azide) |
| **Storage Instruction** | Store at 4°C. |

## Applications

- Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

  Immunohistochemical staining of human liver.

- Western Blot (Cell lysate)

  Western Blot (cell lysate) analysis of HeLa cell lysate.

## Gene Info — HSPD1

| | |
|---|---|
| **Entrez GeneID** | 3329 |
| **Protein Accession#** | P10809 |
| **Gene Name** | HSPD1 |
| **Gene Alias** | CPN60, GROEL, HLD4, HSP60, HSP65, HuCHA60, SPG13 |
| **Gene Description** | heat shock 60kDa protein 1 (chaperonin) |
| **Omim ID** | 118190 605280 |
| **Gene Ontology** | Hyperlink |
| **Gene Summary** | This gene encodes a member of the chaperonin family. The encoded mitochondrial protein may function as a signaling molecule in the innate immune system. This protein is essential for the folding and assembly of newly imported proteins in the mitochondria. This gene is adjacent to a related family member and the region between the 2 genes functions as a bidirectional promoter. Two pseudogenes, both located on chromosome 8, have been associated with this gene. Two transcript variants encoding the same protein have been identified for this gene. Mutations associated with this gene cause autosomal recessive spastic paraplegia 13. [provided by RefSeq |
| **Other Designations** | P60 lymphocyte protein\|chaperonin\|heat shock 60kD protein 1 (chaperonin)\|heat shock protein 65\|mitochondrial heat shock 60kD protein 1 variant 1\|mitochondrial matrix protein P1\|short heat shock protein 60 Hsp60s1\|spastic paraplegia 13 (autosomal dominant) |

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## Pathway

- RNA degradation
- Type I diabetes mellitus

## Disease

- Alzheimer disease
- Arthritis
- Cardiovascular Diseases
- Coronary Disease
- Diabetes Complications
- Diabetes Mellitus
- Disease Progression
- Genetic Predisposition to Disease
- Metabolic Syndrome X
- Multiple Sclerosis
- Neoplasms
- Osteoporosis
- Spastic Paraplegia

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