

## Product datasheet

# Recombinant Human Hsp60 protein ab113177

[2 Images](#)

### Overview

---

<b>Product name</b>	Recombinant Human Hsp60 protein
<b>Protein length</b>	Full length protein

### Description

---

<b>Nature</b>	Recombinant
<b>Source</b>	Escherichia coli

### Amino Acid Sequence

<b>Accession</b>	<a href="#">P10809</a>
<b>Species</b>	Human
<b>Molecular weight</b>	61 kDa
<b>Amino acids</b>	1 to 573

### Specifications

---

Our [Abpromise guarantee](#) covers the use of **ab113177** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

<b>Applications</b>	Western blot Functional Studies SDS-PAGE
<b>Endotoxin level</b>	< 50.000 Eu/mg
<b>Purity</b>	> 90 % SDS-PAGE. ab113177 was purified by multi-step chromatography.
<b>Form</b>	Liquid

### Preparation and Storage

---

<b>Stability and Storage</b>	Shipped on dry ice. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles. Preservative: 0.09% Sodium azide Constituents: 99% PBS, Phosphate Buffer
------------------------------	--

## General Info

---

### Function

Implicated in mitochondrial protein import and macromolecular assembly. May facilitate 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.

### Involvement in disease

Defects in HSPD1 are a cause of spastic paraplegia autosomal dominant type 13 (SPG13) [MIM:605280]. Spastic paraplegia is a degenerative spinal cord disorder characterized by a slow, gradual, progressive weakness and spasticity of the lower limbs.

Defects in HSPD1 are the cause of leukodystrophy hypomyelinating type 4 (HLD4) [MIM:612233]; also called mitochondrial HSP60 chaperonopathy or MitCHAP-60 disease. HLD4 is a severe autosomal recessive hypomyelinating leukodystrophy. Clinically characterized by infantile-onset rotary nystagmus, progressive spastic paraplegia, neurologic regression, motor impairment, profound mental retardation. Death usually occurs within the first two decades of life.

### Sequence similarities

Belongs to the chaperonin (HSP60) family.

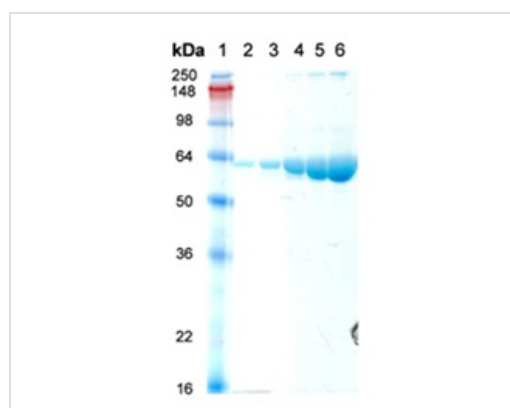
### Cellular localization

Mitochondrion matrix.

---

## Images

---

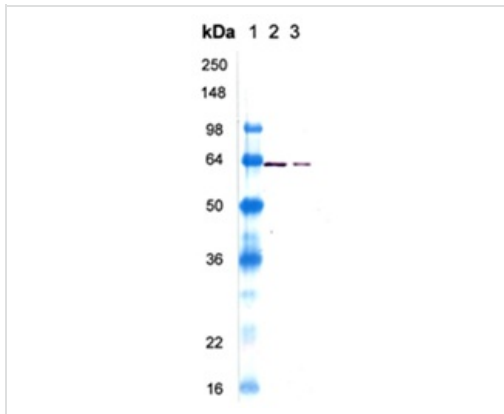


SDS-PAGE analysis of ab113177: Lane 1:

MW marker, Lane 2: 0.5µg, Lane 3: 1µg, Lane

4: 2.5µg, Lane 5: 5µg, Lane 6: 10µg.

SDS-PAGE - Recombinant Human Hsp60 protein  
(ab113177)



Western blot - Recombinant Human Hsp60 protein  
(ab113177)

**All lanes** : a monoclonal anti Hsp60 protein

**Lane 1** : molecular weight marker

**Lane 2** : Recombinant Human Hsp60 protein  
(ab113177) at 0.1  $\mu$ g

**Lane 3** : Recombinant Human Hsp60 protein  
(ab113177) at 0.05  $\mu$ g

**Please note:** All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE"

### Our Abpromise to you: Quality guaranteed and expert technical support

- Replacement or refund for products not performing as stated on the datasheet
- Valid for 12 months from date of delivery
- Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- We investigate all quality concerns to ensure our products perform to the highest standards

If the product does not perform as described on this datasheet, we will offer a refund or replacement. For full details of the Abpromise, please visit <https://www.abcam.com/abpromise> or contact our technical team.

### Terms and conditions

- Guarantee only valid for products bought direct from Abcam or one of our authorized distributors