

## Product datasheet

# Recombinant Human Hsp60 protein ab113177

[1 References](#) [2 Images](#)

### Description

<b>Product name</b>	Recombinant Human Hsp60 protein
<b>Purity</b>	> 90 % SDS-PAGE. ab113177 was purified by multi-step chromatography.
<b>Endotoxin level</b>	< 50.000 Eu/mg
<b>Expression system</b>	Escherichia coli
<b>Accession</b>	<b><u>P10809</u></b>
<b>Protein length</b>	Full length protein
<b>Animal free</b>	No
<b>Nature</b>	Recombinant
<b>Species</b>	Human
<b>Predicted 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>Form</b>	Liquid

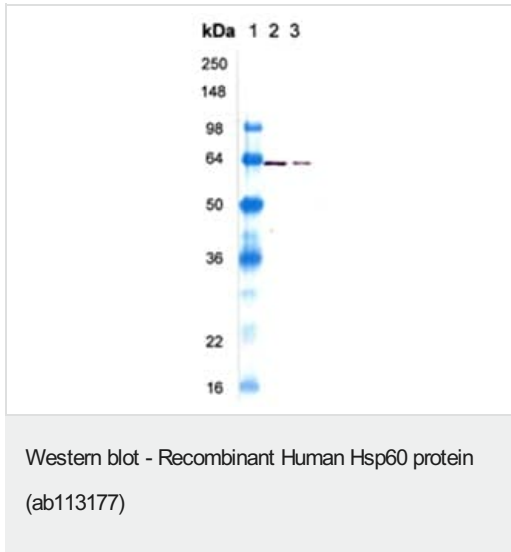
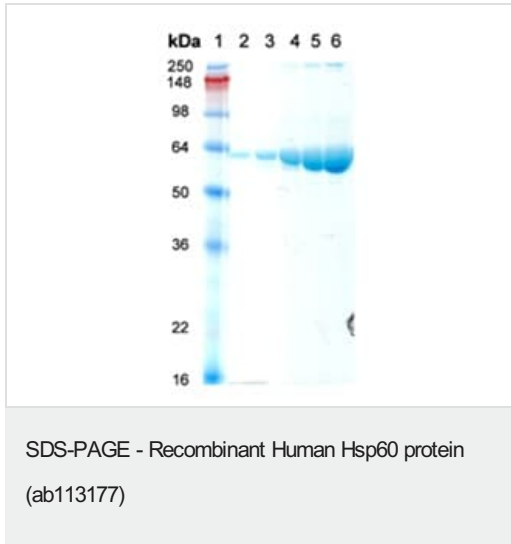
### Preparation and Storage

<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
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### General Info

<b>Function</b>	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.
<b>Involvement in disease</b>	<p>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.</p> <p>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.</p>
<b>Sequence similarities</b>	Belongs to the chaperonin (HSP60) family.
<b>Cellular localization</b>	Mitochondrion matrix.

Images



SDS-PAGE analysis of ab113177: Lane 1: MW marker, Lane 2: 0.5ug, Lane 3: 1ug, Lane 4: 2.5ug, Lane 5: 5ug, Lane 6: 10ug.

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

**Lane 1** : molecualr weight marker

**Lane 2** : Recombinant Human Hsp60 protein (ab113177) at 0.1 µg

**Lane 3** : Recombinant Human Hsp60 protein (ab113177) at 0.05 µg

**Please note:** All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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