

## Product datasheet

# Recombinant Human RPS7 protein ab117224

1 Image

### Overview

<b>Product name</b>	Recombinant Human RPS7 protein
<b>Protein length</b>	Full length protein

### Description

<b>Nature</b>	Recombinant
<b>Source</b>	Escherichia coli
<b>Amino Acid Sequence</b>	
<b>Accession</b>	<a href="#">P62081</a>
<b>Species</b>	Human
<b>Sequence</b>	<p>MGSSHHHHHHSSGLVPRGSHMGSHMFSSSAKIVKPN            GEKPDEFESGISQA            LLELEMNSDLKAQLRELNITAAKEIEVGGGRKAIIFVPV            PQLKSFQKIQ            VRLVRELEKKFSGKHVVVFIQRRILPKPTRKSRTKNKQ            KRPRSRTLAVH            DAILEDLVFPSEVNGKRIRVKLDGSRLIKVHLDKAQQNN            VEHKVETFSGV YKKLTGKDVNFEPPEFQL</p>
<b>Molecular weight</b>	25 kDa including tags
<b>Amino acids</b>	1 to 194
<b>Tags</b>	His tag N-Terminus

### Specifications

Our [Abpromise guarantee](#) covers the use of **ab117224** 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>	<p>SDS-PAGE</p> <p>Mass Spectrometry</p>
<b>Mass spectrometry</b>	MALDI-TOF
<b>Purity</b>	<p>&gt; 90 % SDS-PAGE.</p> <p>ab117224 is purified using conventional chromatography techniques.</p>

**Form** Liquid

## Preparation and Storage

### Stability and Storage

Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -80°C. Avoid freeze / thaw cycle.

pH: 8.00

Constituents: 0.32% Tris HCl, 0.58% Sodium chloride, 30% Glycerol, 0.02% DTT

## General Info

### Function

Required for rRNA maturation.

### Involvement in disease

Defects in RPS7 are the cause of Diamond-Blackfan anemia type 8 (DBA8) [MIM:612563]. DBA8 is a form of Diamond-Blackfan anemia, a congenital non-regenerative hypoplastic anemia that usually presents early in infancy. Diamond-Blackfan anemia is characterized by a moderate to severe macrocytic anemia, erythroblastopenia, and an increased risk of malignancy. 30 to 40% of Diamond-Blackfan anemia patients present with short stature and congenital anomalies, the most frequent being craniofacial (Pierre-Robin syndrome and cleft palate), thumb and urogenital anomalies.

### Sequence similarities

Belongs to the ribosomal protein S7e family.

## Images



15% SDS-PAGE showing ab117224 at approximately 24.7kDa (3µg).

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

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