

## Product datasheet

# Recombinant human Renin protein ab285713

[2 Images](#)

### Description

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<b>Product name</b>	Recombinant human Renin protein
<b>Biological activity</b>	The specific activity is > 2 U/mg. The specific activity of active Renin was assayed using a Renin Activity Fluorometric Assay Kit
<b>Purity</b>	>= 95 % SDS-PAGE.
<b>Expression system</b>	HEK 293 cells
<b>Accession</b>	<b><u>P00797</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>	43 kDa including tags
<b>Tags</b>	His tag C-Terminus
<b>Additional sequence information</b>	8 x His tag

### Specifications

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Our **Abpromise guarantee** covers the use of **ab285713** 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>	SDS-PAGE Functional Studies
<b>Form</b>	Lyophilized
<b>Additional notes</b>	This product is manufactured by BioVision, an Abcam company and was previously called 6300 Human CellExp™ Renin, Human Recombinant. 6300-50 is the same size as the 50 µg size of ab285713.

### Preparation and Storage

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<b>Stability and Storage</b>	Shipped at 4°C. Store at -20°C or -80°C. Avoid freeze / thaw cycle. pH: 8.00
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Constituents: 0.788% Tris HCl, 0.292% Sodium chloride, 5% Trehalose

This product is an active protein and may elicit a biological response in vivo, handle with caution.

## Reconstitution

Centrifuge the vial prior to opening. Reconstitute in sterile PBS (pH 7.4). Do not vortex.

## General Info

### Function

Renin is a highly specific endopeptidase, whose only known function is to generate angiotensin I from angiotensinogen in the plasma, initiating a cascade of reactions that produce an elevation of blood pressure and increased sodium retention by the kidney.

### Involvement in disease

Defects in REN are a cause of renal tubular dysgenesis (RTD) [MIM:267430]. RTD is an autosomal recessive severe disorder of renal tubular development characterized by persistent fetal anuria and perinatal death, probably due to pulmonary hypoplasia from early-onset oligohydramnios (the Potter phenotype).

Defects in REN are the cause of familial juvenile hyperuricemic nephropathy type 2 (HNFJ2) [MIM:613092]. It is a renal disease characterized by juvenile onset of hyperuricemia, slowly progressive renal failure and anemia.

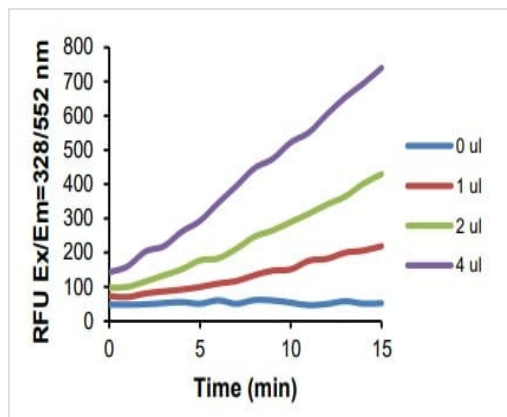
### Sequence similarities

Belongs to the peptidase A1 family.

### Cellular localization

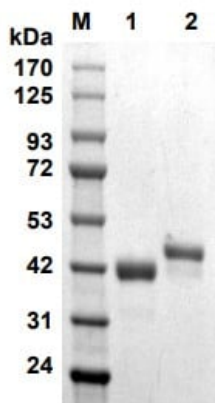
Secreted. Membrane. Associated to membranes via binding to ATP6AP2.

## Images



Renin Enzyme activity: The specific activity of activated Renin is > 2 U/mg assayed by using a Renin Activity Fluorometric Assay Kit. Renin concentration was 0.1 mg/ml

Functional Studies - Recombinant human Renin protein (ab285713)



SDS-PAGE for ab285713 (4-20%) of active and pro Renin: 2 µg of the recombinant Renin and prorenin is loaded under reducing conditions and stained with Coomassie Blue.

Lane M: MW Marker

Lane 1: Active Renin

Lane 2: Prorenin.

SDS-PAGE - Recombinant human Renin protein  
(ab285713)

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

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