

## Product datasheet

# Recombinant human Prorenin protein (Active) ab93266

[3 References](#) [1 Image](#)

### Description

<b>Product name</b>	Recombinant human Prorenin protein (Active)
<b>Biological activity</b>	Fully activatable to renin by catalytic amounts of trypsin. ab93266 is > 95% active.
<b>Purity</b>	> 95 % SDS-PAGE. ab93266 is purified by affinity chromatography.
<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>	44 kDa

### Specifications

Our **Abpromise guarantee** covers the use of **ab93266** 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
<b>Form</b>	Liquid
<b>Additional notes</b>	Extinction coefficient: 1.1 UV Absorbance (280 nm)=0.99

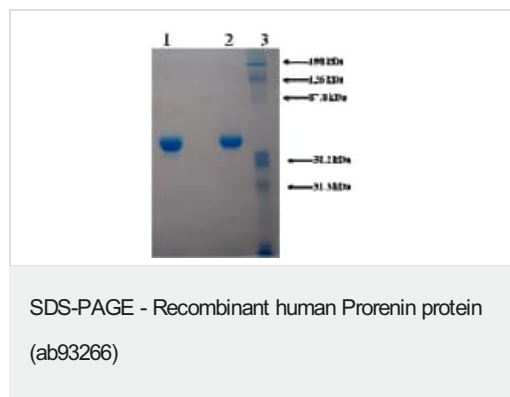
### Preparation and Storage

<b>Stability and Storage</b>	Shipped on dry ice. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles. pH: 8.00 Constituents: 0.79% Tris HCl, 0.29% Sodium chloride This product is an active protein and may elicit a biological response in vivo, handle with caution.
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## General Info

<b>Function</b>	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.
<b>Involvement in disease</b>	<p>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).</p> <p>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.</p>
<b>Sequence similarities</b>	Belongs to the peptidase A1 family.
<b>Cellular localization</b>	Secreted. Membrane. Associated to membranes via binding to ATP6AP2.

## Images



3 ug of ab93266 on 10% SDS-PAGE.

Lane 1: Non-reduced

Lane 2: Reduced

Lane 3: Prestained Standard

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

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