

## Product datasheet

# Recombinant Human Loricrin protein ab114261

1 Image

### Description

<b>Product name</b>	Recombinant Human Loricrin protein
<b>Expression system</b>	Wheat germ
<b>Accession</b>	<a href="#">P23490</a>
<b>Protein length</b>	Full length protein
<b>Animal free</b>	No
<b>Nature</b>	Recombinant
<b>Species</b>	Human
<b>Sequence</b>	<pre> MSYQKKQPTPQPPVDCVKTSGGGGGGGGGGGGCGFF GGGGSGGGSSGSG CGYSGGGGYSGGGCG GGSSGGGGGGGIGGCGGGSGGSVKYSGGGGSSG GGSGCFSSGGGGSGCFSSGGGGSSGGGSGCFS SGGGSSGGGSGCFSS GGGFGSQAVQCQSYGGVSSGGSSGGGSGCFSSGGGG GSVCGYSGGGSG GGSGCGGGSSGGSGSYVSSQQVTQTSCAPQPSYGGG SSGGGGSGGSGC FSSGGGGSSGCGGGG SGIGSGCIISGGGSVCGGGSSGGGGGSSVGG SGSGKGVPICHQTQQKQAPTWPSK </pre>
<b>Predicted molecular weight</b>	61 kDa including tags
<b>Amino acids</b>	1 to 316

### Specifications

Our [Abpromise guarantee](#) covers the use of **ab114261** 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>	ELISA
	SDS-PAGE
	Western blot
<b>Form</b>	Liquid

### Additional notes

## Preparation and Storage

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### Stability and Storage

Shipped on dry ice. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles.

pH: 8.00

Constituents: 0.3% Glutathione, 0.79% Tris HCl

## General Info

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### Function

Major keratinocyte cell envelope protein.

### Involvement in disease

Defects in LOR are a cause of progressive symmetric erythrokeratoderma (PSEK) [MIM:133200]. Erythrokeratodermas are a group of disorders characterized by widespread erythematous plaques, either stationary or migratory, associated with features that include palmoplantar keratoderma. PSEK is characterized by erythematous and hyperkeratotic plaques. Defects in LOR are the cause of Vohwinkel syndrome with ichthyosis (VSI) [MIM:604117]; also known as loricrin keratoderma (LK) or mutilating keratoderma with ichthyosis. VSI is an ichthyotic variant of Vohwinkel syndrome (VS) characterized by progressive symmetric erythrokeratoderma or congenital ichthyosiform erythroderma born as a collodion baby. Common clinical features include hyperkeratosis of the palms and soles with digital constriction.

### Post-translational modifications

Substrate of transglutaminases. Some glutamines and lysines are cross-linked to other loricrin molecules and to SPRRs proteins.

Contains inter- or intramolecular disulfide-bonds.

### Cellular localization

Cytoplasm. Nucleus > nucleoplasm.

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## Images

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ab114261 analysed by 12.5% SDS-PAGE and stained with Coomassie Blue.

SDS-PAGE - Recombinant Human Loricrin protein  
(ab114261)

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

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- Replacement or refund for products not performing as stated on the datasheet
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- 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

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