

## Product datasheet

# Recombinant Human LAMB3 protein ab158811

[1 Image](#)

### Description

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<b>Product name</b>	Recombinant Human LAMB3 protein
<b>Expression system</b>	Wheat germ
<b>Protein length</b>	Protein fragment
<b>Animal free</b>	No
<b>Nature</b>	Recombinant
<b>Species</b>	Human
<b>Sequence</b>	AEGASEQALSAQEGFERIKQKYAELKDRLGQSSMLGEQG ARIQSVKTEAE ELFGETMEMMDRMKDMELELLRGSQAIMLRSADLTGLEK RVEQIRDHING RVLYYATC
<b>Amino acids</b>	1064 to 1171
<b>Tags</b>	GST tag N-Terminus

### Specifications

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Our **Abpromise guarantee** covers the use of **ab158811** 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 ELISA
<b>Form</b>	Liquid

### Additional notes

### Preparation and Storage

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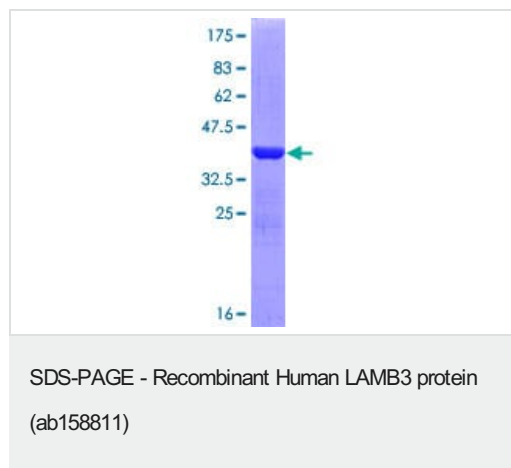
<b>Stability and Storage</b>	Shipped on dry ice. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles. pH: 8.00 Constituents: 0.31% Glutathione, 0.79% Tris HCl
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### General Info

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<b>Function</b>	Binding to cells via a high affinity receptor, laminin is thought to mediate the attachment, migration and organization of cells into tissues during embryonic development by interacting with other extracellular matrix components.
<b>Tissue specificity</b>	Found in the basement membranes (major component).
<b>Involvement in disease</b>	<p>Defects in LAMB3 are a cause of epidermolysis bullosa junctional Herlitz type (H-JEB) [MIM:226700]; also known as junctional epidermolysis bullosa Herlitz-Pearson type. JEB defines a group of blistering skin diseases characterized by tissue separation which occurs within the dermo-epidermal basement membrane. H-JEB is a severe, infantile and lethal form. Death occurs usually within the first six months of life. Occasionally, children survive to teens. H-JEB is marked by bullous lesions at birth and extensive denudation of skin and mucous membranes that may be hemorrhagic.</p> <p>Defects in LAMB3 are a cause of generalized atrophic benign epidermolysis bullosa (GABEB) [MIM:226650]. GABEB is a non-lethal, adult form of junctional epidermolysis bullosa characterized by life-long blistering of the skin, associated with hair and tooth abnormalities.</p>
<b>Sequence similarities</b>	<p>Contains 6 laminin EGF-like domains.</p> <p>Contains 1 laminin N-terminal domain.</p>
<b>Domain</b>	<p>The alpha-helical domains I and II are thought to interact with other laminin chains to form a coiled coil structure.</p> <p>Domain VI is globular.</p>
<b>Cellular localization</b>	Secreted > extracellular space > extracellular matrix > basement membrane.

## Images



ab158811 on a 12.5% SDS-PAGE stained with Coomassie Blue.

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

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