

# *Hev b 10 –His*

(Allergen 10, Superoxiddismutase from *Hevea brasiliensis*, His-tagged)

For research purpose only.



**BIOMAY AG**

Vienna Competence Center  
Lazarettgasse 19 Top 1  
A-1090 Wien

Tel: +43 1 7966296-0  
Fax: +43 1 7966296-111  
e-mail: [info@biomay.com](mailto:info@biomay.com)  
[www.biomay.com](http://www.biomay.com)

## PRODUCT DESCRIPTION:

**Access:** EMBL: AJ249148/Swissprot: Q9STB5

**Mw** = 21,835 Dalton

**Mol. Ext. Coeff.:** 50420; 1mg/ml  $A_{280} = 2.31$

**pI** = 7.9

**Lot#:** 01a

**Amount:** 250 µg

**Quality:** Purity better than 98%

Endotoxin content: 0,125EU/µg

**Reacts with IgE** from Hev b 10 reactive human serum.

---

### General Information:

BIOMAY Hev b 10-His is expressed in *E. coli* as a His-tagged protein. The protein was purified by  $Ni^{2+}$  affinity chromatography and ion exchange chromatography. The product was lyophilized from **1 mM acetic acid**, 2mM  $\beta$ -Mercaptoethanol containing sucrose.

### Reconstitution:

The material can be reconstituted with water or a **weak acidic buffer or solution** (e.g. 1 mM acetic acid). Thorough agitation during dissolution is essential.

If reconstituted to 1 mg/ml the product is soluble to app. 99% and sucrose concentration will be app. 0.3%.

### Storage:

When stored at  $-20^{\circ}\text{C}$  the quality of the material will be maintained for several years. However, for short periods (max. 3 weeks) the lyophilized product may be kept at room temperature. After reconstitution store at  $-20^{\circ}\text{C}$ . Avoid repeated freezing/thawing.

### Quality control:

By SDS-PAGE and staining with Coomassie-Brilliant Blue R-250. Endotoxin content was determined by using a Limulus Amebocyte Lysate (LAL) assay. Immunological properties were controlled by SDS-PAGE/Western-blotting with Hev b 10- specific human IgE.

---

\* The mol.ext.coeff. was calculated from the DNA-derived protein sequence as described by Gill, S.C. and by Hippel, P.H. (1989) Analytical Biochemistry **182**, 319-326.

Please, note that for reasons of shortness only the allergenic content is stated on the product label.

V5. (100121>)

# *Hev b 10 –His (Hev b 10.0102)*

(Allergen 10, Superoxiddismutase from *Hevea brasiliensis*, His-tagged)

For research purpose only.



## BIOMAY AG

Vienna Competence Center  
Lazarettgasse 19 Top 1  
A-1090 Wien

Tel: +43 1 7966296-0  
Fax: +43 1 7966296-111  
e-mail: info@biomay.com  
www.biomay.com

## PRODUCT DESCRIPTION:

**Access:** EMBL: AJ249148/Swissprot: Q9STB5

**Mw** = 21,835 Dalton

**Mol. Ext. Coeff.:** 50420; 1mg/ml  $A_{280} = 2.31$

**pI** = 7.9

**Lot#:** 01

**Amount:** 1 mg

**Quality:** Purity better than 98%

Endotoxin content: 0,125 EU/ $\mu$ g

**Reacts with IgE** from Hev b 10 reactive human serum.

---

### General Information:

BIOMAY Hev b 10-His is expressed in *E. coli* as a His-tagged protein. The protein was purified by  $Ni^{2+}$  affinity chromatography and ion exchange chromatography. The product was lyophilized in 0.1% acetic acid containing 1% sucrose.

### Reconstitution:

The material can be reconstituted with water or a **weak acidic buffer or solution** (e.g 1 mM acetic acid). If reconstituted with water or buffers to 1 mg/ml the product is soluble to app. 99% and sucrose concentration will be app. 0.3%.

Thorough physical suspension of the protein is essential.

### Storage:

The lyophilized product can be kept at room temperature for at least 2 weeks. However, we recommend the product to be stored at  $-20^{\circ}\text{C}$ . Under these conditions the quality of the material will be maintained for several years. The stability at  $4^{\circ}\text{C}$  should at least be 6 months. Reconstituted protein can be stored at  $-20^{\circ}\text{C}$ .

### Quality control:

By SDS-PAGE and staining with Coomassie-blue R250. Endotoxin content was determined by using a Limulus Amebocyte Lysate (LAL) assay. (Immunological properties were controlled by SDS-PAGE\Western-blotting with Hev b 10- specific human IgE)

---

\* The mol.ext.coeff. was calculated from the DNA-derived protein sequence as described by Gill, S.C. and by Hippel, P.H. (1989) Analytical Biochemistry **182**, 319-326.