

# PASTERBACT

Authorised

- Mannheimia haemolytica, Serotype A1, Strain 2806, Leucotoxoid

## Product identification

**Medicine name:**

PASTERBACT

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**Active substance:**

Mannheimia haemolytica, Serotype A1, Strain 2806, Leucotoxoid

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**Target species:**

Sheep

Goat

Cattle (calf)

Sheep (lamb)

Goat (kid)

Cattle (cow)

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**Route of administration:**

Intramuscular use

Subcutaneous use

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## Product details

**Active substance and strength:**

Mannheimia haemolytica, Serotype A1, Strain 2806, Leucotoxoid

20.00 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

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**Pharmaceutical form:**

Emulsion for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Sheep**

- Meat and offal. 0 day

•

**Goat**

- Meat and offal. 0 day

•

**Cattle (calf)**

- Meat and offal. 0 day

•

**Sheep (lamb)**

- Meat and offal. 0 day

•

**Goat (kid)**

- Meat and offal. 0 day

•

**Cattle (cow)**

- Meat and offal. 0 day

**Subcutaneous use:**

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**Sheep**

- Meat and offal. 0 day

•

**Goat**

- Meat and offal. 0 day

- 

**Cattle (calf)**

- Meat and offal. 0 day

- 

**Sheep (lamb)**

- Meat and offal. 0 day

- 

**Goat (kid)**

- Meat and offal. 0 day

- 

**Cattle (cow)**

- Meat and offal. 0 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI02AB04

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Spain

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**Package description:**

Available only in Spanish

Available only in Spanish

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Laboratorios Hipra S.A.

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**Marketing authorisation date:**

1/07/1977

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**Manufacturing sites for batch release:**

Laboratorios Hipra S.A.

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**Responsible authority:**

Spanish Agency For Medicines And Medical Devices

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**Authorisation number:**

2856 ESP

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**Date of authorisation status change:**

18/07/2013

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Labelling

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.