# Histodine

Authorised

- Chlorphenamine maleate

## Product identification

**Medicine name:**
Histodine
Histodine, 10 mg/ml süstelahus veistele

**Active substance:**
Chlorphenamine maleate

**Target species:**
Cattle

**Route of administration:**
Intramuscular use
Intravenous use

## Product details

**Active substance and strength:**
Chlorphenamine maleate
10.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**
Solution for injection

**Withdrawal period by route of administration:**

**Intramuscular use:**

- **Cattle**
  - Milk. 12 hour
  - Meat and offal. 1 day

**Intravenous use:**

- **Cattle**
  - Milk. 12 hour
  - Meat and offal. 1 day

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

QR06AB04

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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:**

Estonia

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**Package description:**

250 ml polypropylene vial, closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box

100 ml polypropylene vial, closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box

250 ml clear Type II glass vial, closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box

100 ml clear Type II glass vial, closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box

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## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Generic application (Article 13(1) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Le Vet. Beheer B.V.

**Marketing authorisation date:**
23/05/2017

**Manufacturing sites for batch release:**
Produlab Pharma B.V.

**Responsible authority:**
State Agency Of Medicines

**Authorisation number:**
2034

**Date of authorisation status change:**
23/05/2017

**Reference member state:**
Netherlands

**Procedure number:**
NL/V/0211/001

**Concerned member states:**
Belgium Cyprus Czechia Estonia France Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Poland Portugal Romania Slovakia Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

## Documents

Summary of Product Characteristics

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

Combined File of all Documents