

Histodine

Authorised

- Chlorphenamine maleate

Product identification

Medicine name:

Histodine

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QR06AB04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Spain

Available in:

Spain

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

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:

8/05/2017

Manufacturing sites for batch release:

Produlab Pharma B.V.

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

3547 ESP

Date of authorisation status change:

26/11/2019

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

Combined File of all Documents

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.

Labelling

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