Histodine 10 mg/ml Solution for Injection for Cattle

Authorised

- Chlorphenamine maleate

Product identification

Medicine name:
Histodine 10 mg/ml Solution for Injection for Cattle

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

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Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

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. B.V.

Marketing authorisation date:
6/07/2017

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

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 41821/4041

Date of authorisation status change:
15/08/2019

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