

PEN-HISTA-STREP SUSPENSION INJECTABLE

Authorised

- Benzylpenicillin procaine monohydrate
- Dexamethasone acetate
- Chlorphenamine maleate
- Dihydrostreptomycin sulfate

Product identification

Medicine name:

PEN-HISTA-STREP SUSPENSION INJECTABLE

Active substance:

Benzylpenicillin procaine monohydrate

Dexamethasone acetate

Chlorphenamine maleate

Dihydrostreptomycin sulfate

Target species:

Cattle

Pig

Cat

Goat

Dog

Route of administration:

Intramuscular use

Intraperitoneal use

Product details

Active substance and strength:

Benzylpenicillin procaine monohydrate

200.69 milligram(s) / 1.00 millilitre(s)

Dexamethasone acetate

0.50 milligram(s) / 1.00 millilitre(s)

Chlorphenamine maleate

9.96 milligram(s) / 1.00 millilitre(s)

Dihydrostreptomycin sulfate

313.05 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Meat and offal. 30 day

- Milk. 6 day

-

Pig

- Meat and offal. 30 day

-

Goat

- Meat and offal. 30 day

- Milk. 6 day

Intraperitoneal use:

-

Cattle

- Meat and offal. 30 day
- Milk. 6 day

-

Pig

- Meat and offal. 30 day

-

Goat

- Meat and offal. 30 day
- Milk. 6 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01RV01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Package description:

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Vetoquinol S.A.

Marketing authorisation date:

2/02/1990

Manufacturing sites for batch release:

Vetoquinol S.A.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/5680265 4/1990

Date of authorisation status change:

2/02/2010

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.

Package Leaflet and Labelling

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