Source URL: https://medicines.health.europa.eu/veterinary/en/600000035563

PENI DHS COOPHAVET SUSPENSION INJECTABLE

Authorised

- Benzylpenicillin procaine monohydrate
- Dihydrostreptomycin sulfate

Product identification

Medicine name:

PENI DHS COOPHAVET SUSPENSION INJECTABLE

Active substance:

Benzylpenicillin procaine monohydrate

Dihydrostreptomycin sulfate

Target species:

Cattle

Pig

Sheep

Goat

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Benzylpenicillin procaine monohydrate 200.70 milligram(s) / 1.00 millilitre(s)

Dihydrostreptomycin sulfate 205.30 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. 7 day

•

Pig

- Meat and offal. 30 day

•

Sheep

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

•

Goat

- Meat and offal. 30 day
- Milk. 7 day

Subcutaneous use:

•

Cattle

- Meat and offal. 30 day
- Milk. 7 day

•

Pig

- Meat and offal. 30 day

•

Sheep

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

•

Goat

- Meat and offal. 30 day
- Milk. 7 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01RA01

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

Available only in $\underline{\mathsf{French}}$

Available only in $\underline{\mathsf{French}}$

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:

Dopharma France S.A.S.

Marketing authorisation date:

6/05/1988

Manufacturing sites for batch release:

Dopharma France S.A.S.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/2943755 1/1988

Date of authorisation status change:

6/05/2013

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.