

OXYTETRIN P SUSPENSION POUR PULVERISATION CUTANEE POUR BOVINS OVINS CAPRINS EQUINS PORCINS ET LAPINS

Authorised

- Oxytetracycline hydrochloride

Product identification

Medicine name:

OXYTETRIN P SUSPENSION POUR PULVERISATION CUTANEE POUR BOVINS OVINS
CAPRINS EQUINS PORCINS ET LAPINS

Active substance:

Oxytetracycline hydrochloride

Target species:

Cattle

Pig

Rabbit

Equid

Sheep

Goat

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Oxytetracycline hydrochloride

107.90 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Cutaneous spray, solution

Withdrawal period by route of administration:

Cutaneous use:

-

Cattle

- Meat and offal. 0 day
- Milk. 0 day

-

Pig

- Meat and offal. 0 day

-

Rabbit

- Meat and offal. 0 day

-

Equid

- Meat and offal. 0 day
- Milk. 0 day

-

Sheep

- Meat and offal. 0 day
- Milk. 0 day

-

Goat

- Meat and offal. 0 day
- Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QD06AA03

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](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Intervet

Marketing authorisation date:

6/04/1988

Manufacturing sites for batch release:

Intervet Productions S.r.l.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/2373469 2/1988

Date of authorisation status change:

6/04/2013

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

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