

OXYTETRA spray

Authorised

- Hydrocortisone acetate
- Oxytetracycline hydrochloride

Product identification

Medicine name:

OXYTETRA spray

Active substance:

Hydrocortisone acetate

Oxytetracycline hydrochloride

Target species:

Cattle

Sheep

Goat

Horse

Pig

Dog

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Hydrocortisone acetate

0.00 gram(s) / 1.00 millilitre(s)

Oxytetracycline hydrochloride

0.02 gram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Cutaneous spray, solution

Withdrawal period by route of administration:

Cutaneous use:

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Cattle

- 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

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Horse

- Meat and offal. 0 day

•

Pig

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QD06AA53

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Bulgaria

Package description:

Available only in [Bulgarian](#)

Available only in [Bulgarian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Bibliographical application (stand-alone)

Marketing authorisation holder:

Farma Vet OOD

Marketing authorisation date:

6/01/2013

Manufacturing sites for batch release:

Farma Vet OOD

Responsible authority:

Bulgarian Food Safety Authority

Authorisation number:

0022-1927

Date of authorisation status change:

6/01/2013

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

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.