Cyclospray 78.6 mg/g, cutaneous spray, suspension for pigs, sheep and cattle

Authorised

- Chlortetracycline hydrochloride

Product identification

Medicine name:
Cyclospray 78.6 mg/g, cutaneous spray, suspension for pigs, sheep and cattle
Cyclospray 78,6 mg/g, δερματικό εκνέφωμα, εναιώρημα για χοίρους, πρόβατα και βοοειδή

Active substance:
Chlortetracycline hydrochloride

Target species:
Cattle
Sheep
Pig

Route of administration:
Cutaneous use

Product details

Active substance and strength:

Chlortetracycline hydrochloride
78.60 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 hour
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD06AA02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Cyprus

Package description:
520 ml (containing 261.52 g) pressurised container of coated tin plate with a plastic valve mechanism and spraying nozzle.

270 ml (containing 130.76 g) pressurised container of coated tin plate with a plastic valve mechanism and spraying nozzle.

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:
Eurovet Animal Health B.V.

Marketing authorisation date:
1/11/2015

Manufacturing sites for batch release:
IGS Aerosols GmbH
Eurovet Animal Health B.V.

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY00530V

Date of authorisation status change:
15/09/2020

Reference member state:
Portugal

Procedure number:
PT/V/0132/001

Concerned member states:
Bulgaria Croatia Cyprus Czechia Estonia Iceland Latvia Romania Slovakia Slovenia Sweden 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

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Summary of Product Characteristics