Animedazon Spray, 2.45 % w/w cutaneous spray suspension for cattle, sheep and pigs

Authorised

- Chlortetracycline hydrochloride

Product identification

Medicine name:
Animedazon Spray, 2.45 % w/w cutaneous spray suspension for cattle, sheep and pigs
ANIMEDAZON 2,45 % w/w dermalno pršilo, suspenzija za govedo, ovce in prašiče

Active substance:
Chlortetracycline hydrochloride

Target species:
Cattle
Sheep
Pig

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Chlortetracycline hydrochloride
3.21 gram(s) / 1.00 Cylinder

Pharmaceutical form:
Cutaneous spray, suspension

Withdrawal period by route of administration:

Cutaneous use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day
- **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:
Slovenia

Available in:
Slovenia

Package description:
(ID2) 12 Cutaneous spray, suspension: unspecified outer container with 12 Cylinder each with 1 Cutaneous spray, suspension

(ID1) 1 Cutaneous spray, suspension: unspecified outer container with 1 Cylinder with 1 Cutaneous spray, suspension

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:
aniMedica GmbH

Marketing authorisation date:
11/08/2015

Manufacturing sites for batch release:
aniMedica GmbH

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
MR/V/0510/001

Date of authorisation status change:
11/08/2015

Reference member state:
Germany

Procedure number:
DE/V/0120/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia France Greece Hungary Iceland Ireland Italy Latvia Lithuania Netherlands Poland Portugal Romania Slovenia Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)
Published on: 14/02/2022
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Package Leaflet

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Labelling

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Combined File of all Documents
