Utertab 2000 mg intrauterine tablet for cattle

Authorised

- Tetracycline hydrochloride

Product identification

Medicine name:
Utertab 2000 mg intrauterine tablet for cattle
Utertab 2 000 mg вътрематочни таблетки за говеда

Active substance:
Tetracycline hydrochloride

Target species:
Cattle (lactating cow)

Route of administration:
Intrauterine use

Product details

Active substance and strength:
Tetracycline hydrochloride
2000.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Intrauterine tablet

Withdrawal period by route of administration:

Intrauterine use:

- **Cattle (lactating cow)**

  - Milk. 96 hour

  - Meat and offal. 10 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG51AA02

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

---

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
(ID9) 500 Intrauterine tablet: unspecified outer container with 100 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine tablet
(ID8) 400 Intrauterine tablet: unspecified outer container with 80 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine tablet
(ID7) 300 Intrauterine tablet: unspecified outer container with 60 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine tablet
(ID6) 200 Intrauterine tablet: unspecified outer container with 40 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine tablet
(ID5) 100 Intrauterine tablet: unspecified outer container with 20 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine tablet
(ID4) 50 Intrauterine tablet: unspecified outer container with 10 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine tablet
(ID3) 20 Intrauterine tablet: unspecified outer container with 4 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine

tablet
(ID2) 10 Intrauterine tablet: unspecified outer container with 2 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine tablet

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Animedica GmbH

Marketing authorisation date:
28/08/2018

Manufacturing sites for batch release:
Animedica GmbH
Animedica Herstellungs GmbH

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2827

Date of authorisation status change:
28/08/2018

Reference member state:
Germany

Procedure number:
DE/V/0176/001

Concerned member states:

Bulgaria Croatia Hungary Ireland Italy Netherlands Poland Portugal Slovakia 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

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.