File downloaded on 2026-11-22
Source URL: https://medicines.health.europa.eu/veterinary/en/600000063544

Utertab 2000 mg intrauterine tablet for cattle

Authorised

- Tetracycline hydrochloride

Product identification

Medicine name:
Utertab 2000 mg intrauterine tablet for cattle

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:

Germany

Available in:

Germany

Package description:

(ID2) 10 Intrauterine tablet: unspecified outer container with 2 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

(ID4) 50 Intrauterine tablet: unspecified outer container with 10 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

(ID6) 200 Intrauterine tablet: unspecified outer container with 40 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
(ID8) 400 Intrauterine tablet: unspecified outer container with 80 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine tablet
(ID9) 500 Intrauterine tablet: unspecified outer container with 100 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:
20/08/2018

Manufacturing sites for batch release:
aniMedica GmbH
aniMedica Herstellungs GmbH

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402466.00.00

Date of authorisation status change:
20/08/2018

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

Combined File of all Documents

English (PDF)
Published on: 25/06/2026
Download

2402466-paren-20181009.rtf