File downloaded on 2026-11-15
Source URL: https://medicines.health.europa.eu/veterinary/en/600000064907

U-tab 2000 mg intrauterine tablet for cattle

Authorised

- Tetracycline hydrochloride

Product identification

Medicine name:
U-tab 2000 mg intrauterine tablet for cattle

Active substance:
Tetracycline hydrochloride

Target species:
Cattle

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**
  - Meat and offal. 10 day
  - Milk. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG51AA02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:

Valid

Authorised in:

Austria

Available in:

Austria

Package description:

(ID4) 10 Intrauterine tablet: Box with 2 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride) each with 5 Intrauterine tablet, closed with Foil (Aluminium)
(ID2) 5 Intrauterine tablet: Box with 1 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride) with 5 Intrauterine tablet, closed with Foil (Aluminium)
(ID3) 20 Intrauterine tablet: Box with 4 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride) each with 5 Intrauterine tablet, closed with Foil (Aluminium)
(ID6) 200 Intrauterine tablet: Box with 40 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride) each with 5 Intrauterine tablet, closed with Foil (Aluminium)
(ID5) 100 Intrauterine tablet: Box with 20 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride) each with 5 Intrauterine tablet, closed with Foil (Aluminium)
(ID1) 50 Intrauterine tablet: Box with 10 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride) each with 5 Intrauterine tablet, closed with Foil (Aluminium)

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:
Eurovet Animal Health B.V.

Marketing authorisation date:
6/04/2011

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

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-00947

Date of authorisation status change:
6/04/2011

Reference member state:
Germany

Procedure number:
DE/V/0140/001

Concerned member states:
Austria Netherlands Poland 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 (RTF)
Published on: 14/03/2026
Download

Labelling

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.