

# U-tab 2000 mg intrauterine tablet for cattle

Authorised

- Tetracycline hydrochloride

## Product identification

**Medicine name:**

U-tab 2000 mg intrauterine tablet for cattle

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**Active substance:**

Tetracycline hydrochloride

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**Target species:**

Cattle

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**Route of administration:**

Intrauterine use

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## Product details

**Active substance and strength:**

Tetracycline hydrochloride  
2000.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Intrauterine tablet

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**Withdrawal period by route of administration:****Intrauterine use:**

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**Cattle**

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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QG51AA02

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Netherlands

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**Available in:**

Netherlands

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**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)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Eurovet Animal Health B.V.

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**Marketing authorisation date:**

20/04/2011

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**Manufacturing sites for batch release:**

Eurovet Animal Health B.V.

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 106878

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**Date of authorisation status change:**

7/02/2022

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0140/001

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**Concerned member states:**

Austria Netherlands Poland United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics