

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:**

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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:

Netherlands

Available in:

Netherlands

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:

20/04/2011

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 106878

Date of authorisation status change:

7/02/2022

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

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