Tetra-Bol 2000 mg tabletes ievadīšanai dzemdē liellopiem

Authorised

- Tetracycline hydrochloride

Product identification

Medicine name:
Tetra-Bol 2000 mg tabletes ievadīšanai dzemdē liellopiem

Active substance:
Tetracycline hydrochloride

Target species:
Cattle (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 (cow)**
  - 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:
Latvia

Package description:
Available only in Latvian
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Dimedium Latvija AS

Marketing authorisation date:
28/10/1995

Manufacturing sites for batch release:
Artesan Pharma GmbH & Co. KG

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/95/0150

Date of authorisation status change:
29/10/1995

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Package Leaflet

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Labelling

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

Summary of Product Characteristics

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