Bovocycline Pessary 2000 mg Intrauterine Tablet for Cattle

Authorised

- Tetracycline hydrochloride

Product identification

Medicine name:
Bovocycline Pessary 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:
United Kingdom (Northern Ireland)

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:
3/06/2011

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

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 16849/4021

Date of authorisation status change:
21/12/2021

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