

Utertab 2000 mg intrauterine tablet for cattle

Authorised

- Tetracycline hydrochloride

Product identification

Medicine name:

Utertab 2000 mg intrauterine tablet for cattle

Utertab 2 000 mg intrauterinná tableta pre hovädzí dobytok

Active substance:

Tetracycline hydrochloride

Target species:

Cattle (lactating 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 (lactating cow)**

- Milk. 96 hour
- Meat and offal. 10 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG51AA02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

Available in:

Slovakia

Package description:

(ID9) 500 Intrauterine tablet: unspecified outer container with 100 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine tablet

(ID8) 400 Intrauterine tablet: unspecified outer container with 80 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine tablet

(ID7) 300 Intrauterine tablet: unspecified outer container with 60 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine tablet

(ID6) 200 Intrauterine tablet: unspecified outer container with 40 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine tablet

(ID5) 100 Intrauterine tablet: unspecified outer container with 20 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine tablet

(ID4) 50 Intrauterine tablet: unspecified outer container with 10 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine

tablet

(ID3) 20 Intrauterine tablet: unspecified outer container with 4 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine tablet

(ID2) 10 Intrauterine tablet: unspecified outer container with 2 Blister (PolyVinyl Chloride; PolyEthylene; PolyVinylidene Chloride; Aluminium) each with 5 Intrauterine tablet

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:

Animedica GmbH

Marketing authorisation date:

29/10/2018

Manufacturing sites for batch release:

Animedica GmbH

Animedica Herstellungs GmbH

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/052/DC/18-S

Date of authorisation status change:

29/10/2018

Reference member state:

Germany

Procedure number:

DE/V/0176/001

Concerned member states:

Bulgaria Croatia Hungary Ireland Italy Netherlands Poland Portugal
Slovakia Spain 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

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