

# Utertab 2000 mg Intrauterine Tablet for Cattle

Authorised

- Tetracycline hydrochloride

## Product identification

**Medicine name:**

Utertab 2000 mg Intrauterine Tablet for Cattle

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**Active substance:**

Tetracycline hydrochloride

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**Target species:**

Cattle (lactating cow)

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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 (lactating cow)**

- Milk. 96 hour
- Meat and offal. 10 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:**

United Kingdom (Northern Ireland)

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**Available in:**

United Kingdom (Northern Ireland)

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**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

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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:**

aniMedica GmbH

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### **Marketing authorisation date:**

21/08/2018

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### **Manufacturing sites for batch release:**

aniMedica GmbH

aniMedica Herstellungs GmbH

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### **Responsible authority:**

The Veterinary Medicines Directorate

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### **Authorisation number:**

Vm 24745/4025

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### **Date of authorisation status change:**

18/06/2024

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### **Reference member state:**

Germany

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**Procedure number:**

DE/V/0176/001

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**Concerned member states:**

Bulgaria Croatia Hungary Ireland Italy Netherlands Poland Portugal  
Slovakia Spain 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

Summary of Product Characteristics

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