

# Gorban 24 % solução injectável destinada a bovinos, equinos, ovinos e suínos

Authorised

- Trimethoprim
- Sulfadoxine

## Product identification

### **Medicine name:**

Gorban 24 % solução injectável destinada a bovinos, equinos, ovinos e suínos

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### **Active substance:**

Trimethoprim

Sulfadoxine

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### **Target species:**

Cattle

Horse

Sheep

Pig

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### **Route of administration:**

Intramuscular use

Subcutaneous use

Intravenous use

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## Product details

### **Active substance and strength:**

Trimethoprim

40.00 milligram(s) / 1.00 millilitre(s)

Sulfadoxine

200.00 milligram(s) / 1.00 millilitre(s)

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### **Pharmaceutical form:**

Solution for injection

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### **Withdrawal period by route of administration:**

#### **Intramuscular use:**

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#### **Cattle**

- Meat and offal. 10 day

- Milk. 4 day

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#### **Horse**

- Meat and offal. 10 day

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#### **Sheep**

- Meat and offal. 15 day

- Milk. 6 day

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#### **Pig**

- Meat and offal. 10 day

#### **Subcutaneous use:**

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#### **Cattle**

- Meat and offal. 10 day

- Milk. 4 day

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**Pig**

- Meat and offal. 10 day

**Intravenous use:**

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**Cattle**

- Meat and offal. 10 day

- Milk. 4 day

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**Pig**

- Meat and offal. 10 day

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**Sheep**

- Meat and offal. 15 day

- Milk. 6 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01EW13

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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:**

Portugal

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**Available in:**

Portugal

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**Package description:**

Available only in Portuguese

Available only in [Portuguese](#)

Available only in [Portuguese](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Virbac

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**Marketing authorisation date:**

22/12/1976

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**Manufacturing sites for batch release:**

Virbac

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**Responsible authority:**

Directorate General For Food And Veterinary

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**Authorisation number:**

651/01/13NFVPT

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**Date of authorisation status change:**

8/01/2019

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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

Combined File of all Documents

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