

# BORGAL 200 mg/ml + 40 mg/ml SOLUCION INYECTABLE

Authorised

- Trimethoprim
- Sulfadoxine

## Product identification

**Medicine name:**

BORGAL 200 mg/ml + 40 mg/ml SOLUCION INYECTABLE

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**Active substance:**

Trimethoprim

Sulfadoxine

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**Target species:**

Cattle

Sheep

Horse

Pig

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**Route of administration:**

Intramuscular 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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**Sheep**

- Meat and offal. 15 day

- Milk. 6 day

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**Horse**

- Meat and offal. 10 day

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**Pig**

- Meat and offal. 10 day

•

**Cattle**

- Meat and offal. 10 day

- Milk. 4 day

•

**Sheep**

- Meat and offal. 15 day

- Milk. 6 day

**Intravenous use:**

- 

**Cattle**

- Meat and offal. 10 day
- Milk. 4 day

- 

**Sheep**

- Meat and offal. 15 day
- Milk. 6 day

- 

**Horse**

- Meat and offal. 10 day

- 

**Cattle**

- Meat and offal. 10 day
- Milk. 4 day

- 

**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:**

Spain

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**Available in:**

Spain

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**Package description:**

Available only in Spanish

Available only in Spanish

Available only in Spanish

Available only in Spanish

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Virbac

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**Marketing authorisation date:**

5/12/2011

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**Manufacturing sites for batch release:**

Virbac

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**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

2416 ESP

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**Date of authorisation status change:**

5/12/2011

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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

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

### Package Leaflet

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

### Labelling

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