

# PENDISTREP 200 mg/ml + 250 mg/ml injection suspension for cattle, sheep, goats and pigs

Authorised

- Dihydrostreptomycin sulfate
- Benzylpenicillin procaine

## Product identification

### Medicine name:

PENDISTREP 200 mg/ml + 250 mg/ml injection suspension for cattle, sheep, goats and pigs

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### Active substance:

Dihydrostreptomycin sulfate  
Benzylpenicillin procaine

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### Target species:

Cattle  
Sheep  
Goat  
Pig  
Horse

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### Route of administration:

Intramuscular use

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## Product details

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

Dihydrostreptomycin sulfate

250.00 milligram(s) / 1.00 millilitre(s)

Benzylpenicillin procaine

200.00 milligram(s) / 1.00 millilitre(s)

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### **Pharmaceutical form:**

Suspension for injection

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

#### **Intramuscular use:**

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#### **Cattle**

- Meat and offal. 28 day

- Milk. 3 day

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#### **Sheep**

- Meat and offal. 28 day

- Milk. 3 day

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#### **Goat**

- Meat and offal. 28 day

- Milk. 3 day

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#### **Pig**

- Meat and offal. 28 day

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#### **Horse**

- Meat and offal. no withdrawal period

Не се разрешава употребата при коне, чието месо е предназначено за консумация от хора.

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

QJ01CE30

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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:**

Bulgaria

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**Available in:**

Bulgaria

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**Package description:**

Available only in [Bulgarian](#)

Available only in [Bulgarian](#)

Available only in [Bulgarian](#)

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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:**

Laboratorios Syva S.A.

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**Marketing authorisation date:**

17/07/2008

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**Manufacturing sites for batch release:**

Laboratorios Syva S.A.

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**Responsible authority:**

Bulgarian Food Safety Authority

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**Authorisation number:**

0022-2097

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**Date of authorisation status change:**

17/07/2008

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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.