

# PROBENCIL 300 mg/ml suspension for injection for cattle and pigs

Authorised

- Benzylpenicillin procaine

## Product identification

**Medicine name:**

PROBENCIL 300 mg/ml suspension for injection for cattle and pigs

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**Active substance:**

Benzylpenicillin procaine

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**Target species:**

Cattle

Pig

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**Route of administration:**

Intramuscular use

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## Product details

**Active substance and strength:**

Benzylpenicillin procaine

300.00 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. 6 day For a treatment duration of 3-5 days
- Meat and offal. 8 day For a treatment duration of 6-7 days
- Milk. 96 hour

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**Pig**

- Meat and offal. 6 day For a treatment duration of 3-5 days
  - Meat and offal. 8 day For a treatment duration of 6-7 days
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01CE09

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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:**

250 ml colourless polyethylene terephthalate (PET) bottle with type I bromobutyl rubber stopper and flip-off cap. Pack size: Carton box with 1 bottle of 250 ml.

100 ml colourless polyethylene terephthalate (PET) bottle with type I bromobutyl rubber stopper and flip-off cap.Pack size: Carton box with 1 vial of 100 ml.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Mevet S.A.

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**Marketing authorisation date:**

7/03/2019

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**Manufacturing sites for batch release:**

Laboratorios Syva S.A.

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**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

3745 ESP

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**Date of authorisation status change:**

7/03/2019

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0398/001

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**Concerned member states:**

Poland Spain

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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

English (PDF)

Published on: 19/01/2025

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### 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.

### Combined File of all Documents

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