

# PermaWay 600 mg intramammary suspension for cattle

Authorised

- Cloxacillin hemibenzathine

## Product identification

**Medicine name:**

PERMAWAY 600 mg intramammary suspension for cattle  
PermaWay 600 mg intramammary suspension for cattle

**Active substance:**

Cloxacillin hemibenzathine

**Target species:**

Cattle (dry cow)

**Route of administration:**

Intramammary use

## Product details

**Active substance and strength:**

Cloxacillin hemibenzathine  
765.40 milligram(s) / 1.00 Syringe

**Pharmaceutical form:**

Intramammary suspension

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**Withdrawal period by route of administration:****Intramammary use:**

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**Cattle (dry cow)**

- Meat and offal. 28 day
- Milk. no withdrawal period

Milk: Interval between treatment and calving is 42 days or longer: 4 days after calving. Interval between treatment and calving is less than 42 days: 46 days after treatment

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

QJ51CF02

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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:**

Ireland

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**Available in:**

Ireland

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**Package description:**

box containing 96 intramammary syringes

box containing 48 intramammary syringes

box containing 24 intramammary syringes

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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:**

Vetoquinol Ireland Limited

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**Marketing authorisation date:**

23/04/2021

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**Manufacturing sites for batch release:**

Vetoquinol Biowet Sp. z o.o.

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA10983/062/001

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**Date of authorisation status change:**

23/04/2021

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0384/001

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**Concerned member states:**

Austria Belgium Cyprus Czechia Estonia France Germany Greece Hungary  
Ireland Italy Latvia Lithuania Luxembourg Malta Netherlands Poland  
Portugal Romania Slovakia Slovenia 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

Combined File of all Documents