

# Parofor 140 mg/ml Solution for use in drinking water/milk

Authorised

- Paromomycin sulfate

## Product identification

**Medicine name:**

Parofor 140 mg/ml Solution for use in drinking water/milk

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**Active substance:**

Paromomycin sulfate

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**Target species:**

Cattle

Pig

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**Route of administration:**

In drinking water/milk use

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## Product details

**Active substance and strength:**

Paromomycin sulfate

200.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for use in drinking water/milk

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**Withdrawal period by route of administration:****In drinking water/milk use:**

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**Cattle**

- Meat and offal. 20 day 20 days for pre-ruminant cattle

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**Pig**

- Meat and offal. 3 day 3 days

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

QA07AA06

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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:**

Malta

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**Package description:**

Parofor 140 mg/ml sol. for drinking water/milk 1000 ml

Parofor 140 mg/ml sol. for drinking water/milk 500 ml

Parofor 140 mg/ml sol. for drinking water/milk 250 ml

Parofor 140 mg/ml sol. for drinking water/milk 125 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application – change in strength (Article 19(1)(a) of Regulation (EU) 2019/6)

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**Marketing authorisation holder:**

HuVepharma

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**Marketing authorisation date:**

17/07/2017

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**Manufacturing sites for batch release:**

Biovet AD

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**Responsible authority:**

Veterinary And Phytosanitary Regulation Department

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**Authorisation number:**

VMA 113

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**Date of authorisation status change:**

17/07/2023

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**Reference member state:**

Belgium

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**Procedure number:**

BE/V/0027/002

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**Concerned member states:**

Austria Bulgaria Cyprus Czechia Denmark Estonia France Germany Greece  
Hungary Ireland Italy Latvia Lithuania Luxembourg Malta Netherlands  
Poland Portugal Romania Slovakia Slovenia Spain  
United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
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## Documents

Combined File of all Documents

English (PDF)

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