

Parofor 175 mg/ml Solution for injection

Not
authorised

- Paromomycin sulfate

Product identification

Medicine name:

Parofor 175 mg/ml Solution for injection

Active substance:

Paromomycin sulfate

Target species:

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Paromomycin sulfate
250.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

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

-

Pig

- Meat and offal. 20 day 20 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA07AA06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Estonia

Package description:

Parofo 175 mg/ml inj. sol. i.m. vial 100 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:

HuVepharma

Marketing authorisation date:

5/11/2018

Manufacturing sites for batch release:

Biovet AD

Responsible authority:

State Agency Of Medicines

Authorisation number:

2119

Date of authorisation status change:

24/02/2025

Reference member state:

Belgium

Procedure number:

BE/V/0027/003

Generic of:

600000086010

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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