

Forespix, 100mg/ml, Solution for injection

Authorised

- Tulathromycin

Product identification

Medicine name:

Forespix, 100mg/ml, Solution for injection

Active substance:

Tulathromycin

Target species:

Cattle

Pig

Sheep

Route of administration:

Subcutaneous use

Intramuscular use

Product details

Active substance and strength:

Tulathromycin

100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

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

-

Cattle

- Meat and offal. 22 day
- Milk. no withdrawal period

Do not use in pregnant animals intended for milk production for human consumption during 2 months prior to expected parturition.,

Intramuscular use:

-

Pig

- Meat and offal. 13 day

-

Sheep

- Meat and offal. 16 day
- Milk. no withdrawal period

Do not use in pregnant animals intended for milk production for human consumption during 2 months prior to expected parturition.,

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA94

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Czechia

Available in:

Czechia

Package description:

Plastic Vial 1 x 50.0 millilitre(s)

Plastic Vial 1 x 100.0 millilitre(s)

Plastic Vial 1 x 250.0 millilitre(s)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Przedsiębiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Marketing authorisation date:

16/12/2020

Manufacturing sites for batch release:

Przedsiębiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Przedsiębiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/077/20-C

Date of authorisation status change:

16/12/2020

Reference member state:

Czechia

Procedure number:

Concerned member states:

Belgium Bulgaria Croatia France Germany Greece Hungary Ireland Italy
Lithuania Netherlands Poland Portugal Romania Slovakia Slovenia Spain

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 5/05/2026

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Package Leaflet

English (PDF)

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Labelling

English (PDF)

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