

# Forespix, 100mg/ml, Solution for injection

Authorised

- Tulathromycin

## Product identification

**Medicine name:**

Forespix, 100mg/ml, Solution for injection

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**Active substance:**

Tulathromycin

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**Target species:**

Cattle

Pig

Sheep

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**Route of administration:**

Subcutaneous use

Intramuscular use

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## Product details

**Active substance and strength:**

Tulathromycin

100.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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

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**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:**

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**Pig**

- Meat and offal. 13 day

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**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.,

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

QJ01FA94

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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:**

Netherlands

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**Available in:**

Netherlands

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**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)

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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:**

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

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**Marketing authorisation date:**

21/01/2021

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**Manufacturing sites for batch release:**

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

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

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 125999

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**Date of authorisation status change:**

29/01/2022

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**Reference member state:**

Czechia

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**Procedure number:**

CZ/V/0166/001

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**Concerned member states:**

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

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

eu-puar-czv0166001-mr-forespix-en.pdf