Forespix, 100mg/ml, Solution for injection

Authorised

- Tulathromycin

Product identification

Medicine name:
Forespix, 100mg/ml, Solution for injection
Forespix, 100 mg/mL, otopina za injekciju, za goveda, svinje i ovce

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:
Croatia

Available in:
Croatia

Package description:
Plastic Vial 1 x 250.0 millilitre(s)
Plastic Vial 1 x 100.0 millilitre(s)
Plastic Vial 1 x 50.0 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:
Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Marketing authorisation date:
9/03/2021

Manufacturing sites for batch release:
Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.
Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/21-01/180

Date of authorisation status change:
22/12/2025

Reference member state:
Czechia

Procedure number:

CZ/V/0166/001

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

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