Florfenikel 300 mg/ml solution for injection for cattle and pigs

Authorised

- Florfenicol

Product identification

Medicine name:
Florfenikel 300 mg/ml solution for injection for cattle and pigs

Active substance:
Florfenicol

Target species:
Cattle
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Florfenicol
300.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 34 day .
  - Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption

- **Pig**
  - Meat and offal. 18 day .

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01BA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

Package description:

Florfenikel 300 mg/ml inj. sol. i.m. vial 250 ml (PP)
Florfenikel 300 mg/ml inj. sol. i.m. vial 100 ml (PP)
Florfenikel 300 mg/ml inj. sol. i.m. vial 12 x 250 ml (PP)
Florfenikel 300 mg/ml inj. sol. i.m. vial 10 x 250 ml (PP)
Florfenikel 300 mg/ml inj. sol. i.m. vial 6 x 250 ml (PP)
Florfenikel 300 mg/ml inj. sol. i.m. vial 12 x 100 ml (PP)
Florfenikel 300 mg/ml inj. sol. i.m. vial 10 x 100 ml (PP)
Florfenikel 300 mg/ml inj. sol. i.m. vial 6 x 100 ml (PP)

Florfenikel 300 mg/ml inj. sol. i.m. vial 6 x 250 ml (glass)
Florfenikel 300 mg/ml inj. sol. i.m. vial 6 x 100 ml (glass)
Florfenikel 300 mg/ml inj. sol. i.m. vial 12 x 250 ml (glass)
Florfenikel 300 mg/ml inj. sol. i.m. vial 10 x 250 ml (glass)
Florfenikel 300 mg/ml inj. sol. i.m. vial 250 ml (glass)
Florfenikel 300 mg/ml inj. sol. i.m. vial 12 x 100 ml (glass)
Florfenikel 300 mg/ml inj. sol. i.m. vial 10 x 100 ml (glass)
Florfenikel 300 mg/ml inj. sol. i.m. vial 100 ml (glass)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
KELA Kempisch Laboratorium Kela Laboratoria

Marketing authorisation date:
2/02/2012

Manufacturing sites for batch release:
S P Veterinaria S.A.
KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:
Danish Medicines Agency

Authorisation number:
47459

Date of authorisation status change:
2/02/2012

Reference member state:
Belgium

Procedure number:
BE/V/0038/001

Concerned member states:
Bulgaria Cyprus Czechia Denmark France Germany Greece Hungary Ireland Italy Luxembourg Netherlands Poland Portugal Romania Slovakia Spain United Kingdom (Northern Ireland)

Generic of:
600000049571

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

Documents

Combined File of all Documents

English (PDF)
Published on: 18/04/2024
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Summary of Product Characteristics

English (PDF)
Published on: 13/03/2026
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Package Leaflet