Zeleris 400 mg/ml florfenicol + 5 mg/ml meloxicam - Solution for injection

Authorised

- Florfenicol
- Meloxicam

Product identification

Medicine name:
Zeleris 400 mg/ml florfenicol + 5 mg/ml meloxicam - Solution for injection

Active substance:
Florfenicol

Meloxicam

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

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

Meloxicam
5.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Meat and offal. 56 day 56 days
  - Milk. no withdrawal period

999 days - Not authorised for use in animals producing milk for human consumption

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01BA99

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain

Package description:

Packaging:Vial (PP/EVOH/PP), Package_size:1 vial, Content:250 ml

Packaging:Vial (PP/EVOH/PP), Package_size:1 vial, Content:100 ml

Packaging:Vial (PP/EVOH/PP), Package_size:1 vial, Content:50 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
15/05/2017

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
15/05/2017

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

Documents

Combined File of all Documents

English (PDF)

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