OVUCRON 25 g/ml

Authorised

- Lecirelin acetate

Product identification

Medicine name:
OVUCRON 25 g/ml
Ovucron, 0,025mg/ml, Injekční roztok

Active substance:
Lecirelin acetate

Target species:
Cattle (cow)
Rabbit (female for reproduction)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Lecirelin acetate
0.03 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Rabbit (female for reproduction)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH01CA92

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Czechia

Package description:
box containing 1 vial of 20 ml
box containing 1 vial of 10 ml
box containing 15 vials of 2 ml

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:
Fatro S.p.A.

Marketing authorisation date:
25/01/2017

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicines

Authorisation number:
96/008/17-C

Date of authorisation status change:
25/01/2017

Reference member state:
Spain

Procedure number:
ES/V/0269/001

Concerned member states:
Czechia Slovakia

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 22/12/2023
Download

Package Leaflet

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Labelling

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