OVUCRON 25 ?g/ml

Authorised

- Lecirelin acetate

Product identification

Medicine name:
OVUCRON 25 ?g/ml
OVUCRON 0,025 mg/ml injekčný roztok pre hovädzí dobytok a králiky

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:
Slovakia

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:
25/05/2017

Manufacturing sites for batch release:
Fatro S.p.A.

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

Authorisation number:
96/020/DC/17-S

Date of authorisation status change:
24/06/2022

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

Combined File of all Documents

eu-PUAR-esv0269001-dcp-ovucron-0-025-mg-ml-en.pdf