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Dalmarelin, 25 mcg/ml süstelahus hobustele, veistele, küülikutele

Authorised

- Lecirelin

Product identification

Medicine name:
Dalmarelin, 25 mcg/ml süstelahus hobustele, veistele, küülikutele

Active substance:
Lecirelin

Target species:
Horse (mare)
Cattle (cow)
Rabbit

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Lecirelin
25.00 microgram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Horse (mare)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Rabbit**
  - 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:

Estonia

Available in:

Estonia

Package description:

Available only in Estonian

Available only in Estonian

Available only in Estonian

Available only in Estonian

Available only in Estonian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
5/12/2002

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

Responsible authority:
State Agency Of Medicines

Authorisation number:
1123

Date of authorisation status change:
5/12/2002

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

Documents

Summary of Product Characteristics

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