

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

This document does not exist in this language (English). You can find it in another language below.