

Luteoplan 0.25 mg/ml solution for injection for cattle and horses

Authorised

- Cloprostenol sodium

Product identification

Medicine name:

Luteoplan 0.25 mg/ml solution for injection for cattle and horses

Luteoplan 0,25 mg/ml oldatos injekció szarvasmarháknak és lovaknak

Active substance:

Cloprostenol sodium

Target species:

Cattle

Horse

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Cloprostenol sodium

0.26 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

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Cattle

- Meat and offal. 1 day
- Milk. 0 hour

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Horse

- Meat and offal. 4 day
- Milk. 24 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG02AD90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Hungary

Package description:

Container size: 20 ml. Container material: Amber Type I glass vials in a cardboard box. Closure: Grey elastomeric bromobutyl rubber stoppers sealed with a plastic flip off button and an aluminium cap.

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:

Syn Vet-Pharma Ireland Limited

Marketing authorisation date:

12/07/2023

Manufacturing sites for batch release:

Laboratorios Ovejero S.A.U.

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

4375/1/23 NÉBIH ÁTI

Date of authorisation status change:

12/07/2023

Reference member state:

Ireland

Procedure number:

IE/V/0771/001

Concerned member states:

Belgium France Germany Hungary Italy Netherlands Poland Portugal Spain
United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to

www.adrreports.eu/vet