Synchromate 250 micrograms/ml solution for injection

Authorised

- Cloprostenol sodium

Product identification

Medicine name:
Synchromate 250 microgramas/ml solução injetável
Synchromate 250 micrograms/ml solution for injection

Active substance:
Cloprostenol sodium

Target species:
Cattle (cow)
Pig (sow)
Horse (mare)
Cattle (heifer)
Pig (sow, nullipar)

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:

- **Cattle (cow)**
  - Meat and offal. 2 day
  - Milk. 0 day
- **Pig (sow)**
  - Meat and offal. 2 day
- **Horse (mare)**
  - Meat and offal. 28 day
- **Cattle (heifer)**
  - Meat and offal. 2 day
  - Milk. 0 day
- **Pig (sow, nullipar)**
  - Meat and offal. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG02AD90

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

Authorisation status:
Valid

Authorised in:
Portugal

Package description:
Cardboard box containing 12 vials of 20ml
Cardboard box containing 5 vials of 20ml
Cardboard box containing 1 vial of 20ml
Cardboard box containing 12 vials of 10ml
Cardboard box containing 5 vials of 10ml
Cardboard box containing 1vial of 10ml
Cardboard box containing 1 vial of 100 ml
Cardboard box containing 1 vial of 50 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:
Alivira Animal Health Limited

Marketing authorisation date:
27/09/2022

Manufacturing sites for batch release:
Bremer Pharma GmbH

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
1532/01/22DFVPT

Date of authorisation status change:
7/10/2025

Reference member state:
Spain

Procedure number:
ES/V/0411/001

Concerned member states:
Belgium Hungary Italy Netherlands Poland Portugal Romania

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

Documents

Summary of Product Characteristics

Package Leaflet

Labelling

Combined File of all Documents

English (PDF)
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