

Synchromate 0.25 mg/ml solution for injection for cattle, pigs and horses

Authorised

- Cloprostenol sodium

Product identification

Medicine name:

Synchromate 0.25 mg/ml solution for injection for cattle, pigs and horses

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
263.00 microgram(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:

Hungary

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 1 vial 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:

24/10/2022

Manufacturing sites for batch release:

Bremer Pharma GmbH

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

4342/X/22 NÉBIH ÁTI (1 db 10 ml-es injekciós üvege)

Date of authorisation status change:

24/10/2022

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

eu-PUAR-synchromate-0.25mg-ml--solution-for-injection-for-cattle--pigs-and-horses-
en.pdf