VETEGLAN 0.075 mg/ml Solution for injection for cows, sows and mares

Authorised

- Cloprostenol sodium

Product identification

Medicine name:
VETEGLAN 0.075 mg/ml Solution for injection for cows, sows and mares
VETEGLAN 0.075 mg/ml инжекционен разтвор за крави, свине и кобили

Active substance:
Cloprostenol sodium

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

Route of administration:
Solution for injection

Product details

Active substance and strength:
Cloprostenol sodium
0.08 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Solution for injection:

- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Pig**
  - Meat and offal. 1 day
- **Horse (mare)**
  - Meat and offal. 2 day
  - Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG02AD90

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
20 ml amber coloured Type I glass vials, with Teflon-coated chlorobutyl rubber closures and aluminium seals with blue coloured plastic flip-offs, packaged singly in a cardboard box.
10 ml amber coloured Type I glass vials, with Teflon-coated chlorobutyl rubber closures and aluminium seals with blue coloured plastic flip-offs, packaged singly in a cardboard box.

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:
Laboratorios Calier S.A.

Marketing authorisation date:
24/01/2017

Manufacturing sites for batch release:
Laboratorios Calier S.A.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2706

Date of authorisation status change:
16/01/2023

Reference member state:
Portugal

Procedure number:
PT/V/100/001

Concerned member states:
Austria Belgium Bulgaria Croatia Denmark France Germany Greece Hungary Ireland Latvia Lithuania Netherlands Romania Spain United Kingdom (Northern Ireland)

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.

Package Leaflet

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

Labelling

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