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

Active substance:
Cloprostenol sodium

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

Route of administration:
Intramuscular use

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:
Intramuscular use:

- **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:
Austria

Package description:
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.
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.

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:
27/04/2017

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

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
837610

Date of authorisation status change:
27/04/2017

Reference member state:
Portugal

Procedure number:
PT/V/0100/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus 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

Labelling

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.

Summary of Product Characteristics

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

Combined File of all Documents