PGF Veyx 0.0875 mg/ml solution for injection for cattle and pigs

Authorised

- Cloprostenol sodium

Product identification

Medicine name:
PGF Veyx 0.0875 mg/ml solution for injection for cattle and pigs

Active substance:
Cloprostenol sodium

Target species:
Cattle (heifer)
Pig (sow)
Cattle (cow)

Route of administration:
Intramuscular use

Product details

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

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle (heifer)**
  - Milk. 0 hour
  - Meat and offal. 1 day
- **Pig (sow)**
  - Meat and offal. 1 day
- **Cattle (cow)**
  - Meat and offal. 1 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:

Germany

Available in:

Germany

Package description:

(ID1) 10 millilitre(s): unspecified outer container with 1 Vial (glass) with 10 millilitre(s)
(ID2) 20 millilitre(s): unspecified outer container with 1 Vial (glass) with 20 millilitre(s)
(ID3) 50 millilitre(s): unspecified outer container with 1 Vial (glass) with 50 millilitre(s)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Veyx Pharma GmbH

Marketing authorisation date:
21/05/2012

Manufacturing sites for batch release:
Veyx Pharma GmbH
Veyx-Pharma B.V.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401538.00.00

Date of authorisation status change:
14/09/2017

Reference member state:
Germany

Procedure number:
DE/V/0146/001

Concerned member states:
Austria Belgium Bulgaria Czechia Estonia France Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)
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