PGF Veyx forte 0.250 mg/ml solution for injection for cattle and pigs

Authorised

- Cloprostenol sodium

Product identification

Medicine name:
PGF Veyx forte 0.250 mg/ml solution for injection for cattle and pigs
PGF VEYX

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

0.26 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)**
  - Milk. 0 hour
  - Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG02AD90

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

Authorisation status:
Valid

Authorised in:
Italy

Available in:
Italy

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

(ID1) 10 millilitre(s): unspecified outer container with 1 Vial (glass) with 10 millilitre(s)

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:
Veyx Pharma GmbH

Marketing authorisation date:
11/12/2012

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

Responsible authority:
Ministry Of Health

Authorisation number:
104409

Date of authorisation status change:
11/12/2012

Reference member state:
Germany

Procedure number:
DE/V/0146/002

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

2401539-paren-20131030.pdf