Cyclix 250 microgram/ml solution for injection for cattle

Authorised

- Cloprostenol sodium

Product identification

Medicine name:
Cyclix 250 microgram/ml solution for injection for cattle
Cyclix 250 mikrogram/ml injektionsvæske, opløsning

Active substance:
Cloprostenol sodium

Target species:
Cattle (cow)

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. 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:
Denmark

Available in:
Denmark

Package description:
(ID2) 50 millilitre(s): unspecified outer container with 1 Vial (Glass) with 50 millilitre(s)
(ID1) 20 millilitre(s): unspecified outer container with 1 Vial (Glass) with 20 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:
Virbac

Marketing authorisation date:
10/03/2006

Manufacturing sites for batch release:
Virbac

Responsible authority:
Danish Medicines Agency

Authorisation number:
38339

Date of authorisation status change:
10/03/2006

Reference member state:
Germany

Procedure number:
DE/V/0111/001

Concerned member states:
Austria Belgium Czechia Denmark Finland Greece Hungary Ireland Italy Luxembourg Netherlands Poland Portugal Slovakia Slovenia 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

English (PDF)
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