

CYCLIX

Authorised

- Cloprostenol sodium

Product identification

Medicine name:

CYCLIX

Active substance:

Cloprostenol sodium

Target species:

Cattle (cow for reproduction)

Cattle (heifer)

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:

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Cattle (cow for reproduction)

- Meat and offal. 2 day
- Milk. 0 day

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Cattle (heifer)

- Meat and offal. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG02AD90

Legal status of supply:

This information is not available for this product.

Authorisation status:

Valid

Authorised in:

Romania

Available in:

Romania

Package description:

Available only in Romanian

Available only in Romanian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:

Virbac

Marketing authorisation date:

5/06/2007

Manufacturing sites for batch release:

Virbac

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

160267

Date of authorisation status change:

22/01/2024

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.