

Cyclofin 300 mg/ml + 20 mg/ml solution for injection for cattle

Authorised

- Flunixin
- Oxytetracycline

Product identification

Medicine name:

Cyclofin 300 mg/ml + 20 mg/ml Injektionslösung für Rinder
Cyclofin 300 mg/ml + 20 mg/ml solution for injection for cattle

Active substance:

Flunixin
Oxytetracycline

Target species:

Cattle

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Flunixin
20.00 milligram(s) / 1.00 millilitre(s)
Oxytetracycline
300.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intramuscular use:****. Cattle**

- Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01AA56

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria

Package description:

Type II, clear glass vial of 100 ml, with a 20 mm bromobutyl rubber stopper, and aluminium cap. One glass vial is packaged in a cardboard box.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:

Dechra Regulatory B.V.

Marketing authorisation date:

11/08/2023

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

841750

Date of authorisation status change:

11/08/2023

Reference member state:

Ireland

Procedure number:

IE/V/0773/001

Concerned member states:

Austria Belgium France Germany Italy Netherlands Poland Portugal 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)

Published on: 3/05/2024

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Labelling

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Package Leaflet

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