PermaWay 600 mg intramammary suspension for cattle

Authorised

- Cloxacillin hemibenzathine

Product identification

Medicine name:
PermaWay 600 mg intramammary suspension for cattle
Permaway 600 mg Suspensie voor intramammair gebruik
Permaway 600 mg Suspension intramammaire
Permaway 600 mg Suspension zur intramammären Anwendung

Active substance:
Cloxacillin hemibenzathine

Target species:
Cattle (dry cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Cloxacillin hemibenzathine

765.40 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:
Intramammary use:

- **Cattle (dry cow)**
  - Meat and offal. 28 day
  - Milk. no withdrawal period

Milk: Interval between treatment and calving is 42 days or longer: 4 days after calving.Interval between treatment and calving is less than 42 days: 46 days after treatment

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51CF02

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

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
box containing 96 intramammary syringes
box containing 48 intramammary syringes
box containing 24 intramammary syringes

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:
Vetoquinol

Marketing authorisation date:
31/08/2021

Manufacturing sites for batch release:
Vetoquinol Biowet Sp. z o.o.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V589635

Date of authorisation status change:
8/10/2021

Reference member state:
Spain

Procedure number:
ES/V/0384/001

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

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

Documents

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

English (PDF)
Published on: 22/12/2023
Download

Combined File of all Documents

Labelling

This document does not exist in this language (English). You can find it in another language below.