Ubroseal blue Dry Cow 2.6 g intramammary suspension for cattle

Authorised

- Bismuth subnitrate, heavy

Product identification

Medicine name:
Ubroseal blue Dry Cow 2.6 g intramammary suspension for cattle
Ubroxeal blue Dry Cow 2.6 g Suspensie voor intramammair gebruik
Ubroxeal blue Dry Cow 2.6 g Suspension intramammaire
Ubroxeal blue Dry Cow 2.6 g Suspension zur intramammären Anwendung

Active substance:
Bismuth subnitrate, heavy

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
Bismuth subnitrate, heavy
2.60 gram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG52X

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

Authorisation status:
Valid

Authorised in:
Belgium

Package description:
A 4g polyethylene intramammary syringe consisting of a barrel with plunger and a polyethylene dual-cap.Cardboard box of 20 syringes and 20 cleaning towels
A 4g polyethylene intramammary syringe consisting of a barrel with plunger and a polyethylene dual-cap.Polyethylene bucket of 60 syringes and 60 cleaning towels
A 4g polyethylene intramammary syringe consisting of a barrel with plunger and a polyethylene dual-cap.Polyethylene bucket of 120 syringes and 120 cleaning towels

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:

Univet Limited

Marketing authorisation date:
16/01/2018

Manufacturing sites for batch release:
Univet Limited

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V521955

Date of authorisation status change:
10/02/2021

Reference member state:
Ireland

Procedure number:
IE/V/0437/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Italy Latvia Liechtenstein Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden 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: 11/02/2022
Download

Package Leaflet

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