Ubroseal blue Dry Cow 2.6 g intramammary suspension for cattle

Authorised

- Bismuth subnitrate, heavy

Product identification

Medicine name:
Ubroxeal blue Dry Cow 2,6 g suspension intramammaire pour bovins
Ubroxeal blue Dry Cow 2,6 g Suspension zur intramammären Anwendung bei Rindern
Ubroseal blue Dry Cow 2.6 g intramammary suspension for cattle

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:
Luxembourg

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:
Ministry Of Health And Social Security

Authorisation number:
V 721/18/02/1656

Date of authorisation status change:
16/01/2018

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
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Package Leaflet

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