Noroseal 2.6 g Intramammary Suspension for Cattle

Not authorised

- Bismuth subnitrate, heavy

Product identification

Medicine name:
Noroseal 2.6 g Intramammary Suspension for Cattle
Intraseal 2.6 g Suspensie voor intramammair gebruik
Intraseal 2.6 g Suspension intramammaire
Intraseal 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

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Authorisation status:
Surrendered

Authorised in:
Belgium

Package description:
Low density polyethylene syringe with a smooth tapered hermetically sealed nozzle.
Pack sizes: Buckets of 120 syringes including 120 individually wrapped teat cleaning towels.
Low density polyethylene syringe with a smooth tapered hermetically sealed nozzle.
Pack sizes: Cartons of 60 syringes including 60 individually wrapped teat cleaning towels.
Low density polyethylene syringe with a smooth tapered hermetically sealed nozzle.
Pack sizes: Cartons of 24 syringes including 24 individually wrapped teat 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:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
2/09/2013

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V442321

Date of authorisation status change:
14/06/2023

Reference member state:
Ireland

Procedure number:
IE/V/0587/001

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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Package Leaflet

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