Sureseal 2.6 g Intramammary Suspension for Cattle

Authorised

- Bismuth subnitrate, heavy

Product identification

Medicine name:
Sureseal 2.6 g Intramammary Suspension for Cattle
Intraseal 2,6 g suspensão intramamária para bovinos (vacas leiteiras).

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 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG52X

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

Authorisation status:
Valid

Authorised in:
Portugal

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:
4/07/2013

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

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
686/01/13DFVPT

Date of authorisation status change:
5/12/2024

Reference member state:
Ireland

Procedure number:
IE/V/0586/001

Concerned member states:
Belgium France Italy Netherlands 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

Combined File of all Documents

English (PDF)
Published on: 26/01/2025
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