Noroseal 2.6 g Intramammary Suspension for Cattle

Authorised

- Bismuth subnitrate, heavy

Product identification

Medicine name:
Noroseal 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

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Authorisation status:
Valid

Authorised in:
Sweden

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:
29/08/2013

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

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
48339

Date of authorisation status change:
29/08/2013

Reference member state:
Ireland

Procedure number:
IE/V/0587/001

Concerned member states:
Austria Belgium Bulgaria Czechia Denmark Estonia Finland France Germany Greece Hungary Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia Spain Sweden

United Kingdom (Northern Ireland)

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

Documents

Package Leaflet

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Summary of Product Characteristics

English (PDF)
Published on: 2/03/2025
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Labelling

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Combined File of all Documents