

# Noroseal 2.6 g Intramammary Suspension for Cattle

Authorised

- Bismuth subnitrate, heavy

## Product identification

**Medicine name:**

Noroseal 2.6 g Intramammary Suspension for Cattle

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**Active substance:**

Bismuth subnitrate, heavy

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**Target species:**

Cattle

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**Route of administration:**

Intramammary use

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## Product details

**Active substance and strength:**

Bismuth subnitrate, heavy  
2.60 gram(s) / 1.00 Syringe

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**Pharmaceutical form:**

Intramammary suspension

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**Withdrawal period by route of administration:****Intramammary use:**

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**Cattle**

- Meat and offal. 0 day

- Milk. 0 hour

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QG52X

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**Legal status of supply:**

Veterinary medicinal product not subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Estonia

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**Available in:**

Estonia

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Norbrook Laboratories (Ireland) Limited

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**Marketing authorisation date:**

26/09/2013

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**Manufacturing sites for batch release:**

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

Norbrook Laboratories Limited

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**Responsible authority:**

State Agency Of Medicines

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**Authorisation number:**

1786

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**Date of authorisation status change:**

26/09/2013

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0587/001

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**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)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

English (PDF)

Published on: 2/03/2025

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Combined File of all Documents