Sureseal 2.6 g Intramammary Suspension for Cattle

Authorised

- Bismuth subnitrate, heavy

Product identification

Medicine name:
Sureseal 2.6 g Intramammary Suspension for Cattle
NOROSEAL 2,6 g SUSPENSION INTRAMAMARIA PARA BOVINO

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:
Spain

Available in:
Spain

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:
8/08/2013

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

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
2875 ESP

Date of authorisation status change:
26/02/2018

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

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

Labelling

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents