ORBESEAL, 2.6g, Intramamární suspenze

Authorised

- Bismuth subnitrate

Product identification

Medicine name:
ORBESEAL, 2.6g, Intramamární suspenze

Active substance:
Bismuth subnitrate

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
Bismuth subnitrate
2.60 gram(s) / 1.00 Applicator

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

Authorisation status:
Valid

Authorised in:
Czechia

Available in:
Czechia

Package description:
Available only in Czech
Available only in Czech
Available only in Czech

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Ceska Republika s.r.o.

Marketing authorisation date:
10/04/2004

Manufacturing sites for batch release:
Haupt Pharma Latina S.r.l.
CROSS VETPHARM GROUP Ltd.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/021/04-C

Date of authorisation status change:
10/04/2004

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

Documents

Summary of Product Characteristics

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Package Leaflet

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

Labelling

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