

ORBESEAL DRY COW 2.6G INTRAMAMMARY SUSPENSION

Authorised

- Bismuth subnitrate, heavy

Product identification

Medicine name:

ORBESEAL DRY COW 2.6G INTRAMAMMARY SUSPENSION

Orbeseal Dry Cow, 2,6 g, intramamarna suspenzija za goveda

Active substance:

Bismuth subnitrate, heavy

Target species:

Cattle (dry cow)

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:**

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Cattle (dry cow)

- 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:

Croatia

Package description:

Boxe of 24 syringes

Plastic bucket of 120 syringes

Boxes of 60 syringes

Plastic bucket of 144 syringes

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Zoetis B.V.

Marketing authorisation date:

25/09/2014

Manufacturing sites for batch release:

CROSS VETPHARM GROUP Ltd.

Haupt Pharma Latina S.r.l.

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/19-01/78

Date of authorisation status change:

27/08/2024

Reference member state:

France

Procedure number:

FR/V/0341/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Denmark Finland Germany Greece
Ireland Italy Luxembourg Netherlands Norway Portugal Romania Slovenia
Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

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