ORBESEAL DRY COW 2.6G INTRAMAMMARY SUSPENSION

Authorised

- Bismuth subnitrate, heavy

Product identification

Medicine name:
ORBESEAL DRY COW 2.6G INTRAMAMMARY SUSPENSION
ORBESEAL

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:

- **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:
Italy

Available in:
Italy

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 Italia S.r.l.

Marketing authorisation date:
10/06/2003

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

Responsible authority:
Ministry Of Health

Authorisation number:
103593

Date of authorisation status change:
10/06/2003

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

Combined File of all Documents

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