Cepravin Dry Cow 250 mg Intramammary suspension

Authorised

- Cefalonium dihydrate

Product identification

Medicine name:
Cepravin Dry Cow 250 mg Intramammary suspension

Active substance:
Cefalonium dihydrate

Target species:
Cattle (dairy cow at drying-off)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Cefalonium dihydrate
269.60 milligram(s) / 1.00 Applicator

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle (dairy cow at drying-off)**
  - Meat and offal. 21 day
  - Milk. 96 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51DB90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
(ID1) 60 gram(s): Box (cardboard) with 20 Applicator (polyethylene) each with 3 gram(s), closed with Lid (polyethylene)
(ID2) 432 gram(s): container with 144 Applicator (polyethylene) each with 3 gram(s), closed with Lid (polyethylene)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet Nederland B.V.

Marketing authorisation date:

28/03/2013

Manufacturing sites for batch release:
Trirx Segre
Intervet International GmbH

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 110469

Date of authorisation status change:
28/03/2013

Reference member state:
Germany

Procedure number:
DE/V/0184/001

Concerned member states:
Austria Bulgaria Cyprus Estonia Latvia Netherlands

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

Documents

Combined File of all Documents

English (PDF)
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