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RILEXINE DC 375 MG INTRAMAMMARY SUSPENSION FOR DRY COWS

Authorised

- Cephalexin benzathine

Product identification

Medicine name:
RILEXINE DC 375 MG INTRAMAMMARY SUSPENSION FOR DRY COWS

Active substance:
Cephalexin benzathine

Target species:
Cattle (dry cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Cephalexin benzathine
500.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle (dry cow)**
  - Meat and offal. 4 day
  - Milk. 43 day 42.5 days after treatment when dry period is 42 days or less
  - Milk. 12 hour 12 hours after calving when dry period is more than 42 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51DB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Germany

Available in:

Germany

Package description:

Box of 12 x 8g intramammary syringes and 12 cleaning towels
Box of 60 x 8g intramammary syringes and 60 cleaning towels
Box of 24 x 8g intramammary syringes and 24 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:
Virbac

Marketing authorisation date:
22/03/2022

Manufacturing sites for batch release:
Haupt Pharma Latina S.r.l.
Virbac

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
V7004231.00.00

Date of authorisation status change:
22/03/2022

Reference member state:
France

Procedure number:
FR/V/0438/001

Concerned member states:
Austria Belgium Bulgaria Croatia Czechia Denmark Estonia Finland Germany Hungary Ireland Italy Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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Documents

Combined File of all Documents

English (PDF)
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