Rilexine 200 LC

Authorised

- Cefalexin monohydrate

Product identification

Medicine name:
Rilexine 200 LC

Active substance:
Cefalexin monohydrate

Target species:
Cattle (dairy cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Cefalexin monohydrate
210.40 milligram(s) / 1.00 Applicator

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle (dairy cow)**
  - Meat and offal. 4 day
  - Milk. 3 day

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:
Available only in German
Available only in German
Available only in German

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Virbac

Marketing authorisation date:

26/09/2000

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

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
400144.00.00

Date of authorisation status change:
7/11/2005

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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