Rilexine 200 T intramammary suspension for cattle

Authorised

- Cefalexin

Product identification

Medicine name:
Rilexine 200 T intramammary suspension for cattle

Active substance:
Cefalexin

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
Cefalexin
200.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle**
  - Meat and offal. 4 day
  - Milk. 48 hour Мляко: 48 часа (4 издоявания)

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51DB01

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian

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:
Virbac

Marketing authorisation date:
8/02/2010

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

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2534

Date of authorisation status change:
8/02/2010

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