

Rilexine 200 suspensija ievadīšanai tesmenī liellopiem

Authorised

- Cefalexin

Product identification

Medicine name:

Rilexine 200 suspensija ievadīšanai tesmenī liellopiem

Active substance:

Cefalexin

Target species:

Cattle (lactating cow)

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:**

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Cattle (lactating cow)

- Milk. 2 day 2 dienas (4 slaukšanas reizes)

- Meat and offal. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51DB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Latvia

Package description:Available only in LatvianAvailable only in Latvian

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:

27/09/2007

Manufacturing sites for batch release:

Virbac

Haupt Pharma Latina S.r.l.

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/NRP/07/1702

Date of authorisation status change:

27/09/2007

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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