

# RILEXINE 200 intramamálna suspenzia

Authorised

- Cefalexin monohydrate

## Product identification

**Medicine name:**

RILEXINE 200 intramamálna suspenzia

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**Active substance:**

Cefalexin monohydrate

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**Target species:**

Cattle (lactating cow)

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**Route of administration:**

Intramammary use

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## Product details

**Active substance and strength:**

Cefalexin monohydrate  
210.40 milligram(s) / 1.00 Syringe

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**Pharmaceutical form:**

Intramammary suspension

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**Withdrawal period by route of administration:**

**Intramammary use:**

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**Cattle (lactating cow)**

- Meat and offal. 4 day

- Milk. 48 hour  
4 milkings

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ51DB01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Slovakia

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**Available in:**

Slovakia

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**Package description:**

Available only in [Slovak](#)

Available only in [Slovak](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Virbac

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**Marketing authorisation date:**

13/05/1993

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**Manufacturing sites for batch release:**

Haupt Pharma Latina S.r.l.

Virbac

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

96/046/93-S

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**Date of authorisation status change:**

13/05/1993

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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