

# Novocillin LC 1000 mg intramammary suspension for lactating cow

Authorised

- Oxacillin sodium monohydrate

## Product identification

**Medicine name:**

Novocillin LC 1000 mg intramammary suspension for lactating cow

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**Active substance:**

Oxacillin sodium monohydrate

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**Target species:**

Cattle (dairy cow)

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**Route of administration:**

Intramammary use

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## Product details

**Active substance and strength:**

Oxacillin sodium monohydrate  
1042.50 milligram(s) / 1.00 Applicator

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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 (dairy cow)**

- Meat and offal. 6 day
  - Milk. 144 hour
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ51CF04

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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:**

Germany

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**Available in:**

Germany

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**Package description:**

(ID3) 240 gram(s): Box (cardboard) with 24 Applicator (linear low-density polyethylene) each with 10 gram(s)

(ID2) 200 gram(s): Box (cardboard) with 20 Applicator (linear low-density polyethylene) each with 10 gram(s)

(ID1) 100 gram(s): Box (cardboard) with 10 Applicator (linear low-density polyethylene) each with 10 gram(s)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Pharmanovo Veterinaerarztneimittel GmbH

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**Marketing authorisation date:**

9/12/2020

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**Manufacturing sites for batch release:**

Vet-Agro Trading Sp. z o.o.

Produlab Pharma B.V.

Przedsiębiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

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**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

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**Authorisation number:**

402684.00.00

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**Date of authorisation status change:**

9/12/2020

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0333/001

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**Concerned member states:**

Austria Bulgaria Cyprus Czechia Estonia France Hungary Iceland Ireland

Italy Lithuania Poland Portugal Romania Spain

United Kingdom (Northern Ireland)

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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

English (PDF)

Published on: 19/03/2025

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