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# Carepen vet 600 mg intramammary suspension for lactating cows

Authorised

- Benzylpenicillin (procaine) monohydrate

## Product identification

**Medicine name:**
Carepen vet 600 mg intramammary suspension for lactating cows

**Active substance:**
Benzylpenicillin (procaine) monohydrate

**Target species:**
Cattle (lactating cow)

**Route of administration:**
Intramammary use

## Product details

**Active substance and strength:**
Benzylpenicillin (procaine) monohydrate
600.00 milligram(s) / 1.00 Syringe

**Pharmaceutical form:**

Intramammary suspension

**Withdrawal period by route of administration:**

**Intramammary use:**

- **Cattle (lactating cow)**
  - Meat and offal. 3 day
  - Milk. 6 day

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

QJ51CE09

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**

Valid

**Authorised in:**

Norway

**Package description:**

Pack size: 100 x 10 g with 100 cleaning towels. White intramammary syringe (LDPE) with a double tip (LDPE) packed in a cardboard container.
Pack size: 40 x 10 g with 40 cleaning towels. White intramammary syringe (LDPE) with a double tip (LDPE) packed in a cardboard container.
Pack size: 20 x 10 g with 20 cleaning towels. White intramammary syringe (LDPE) with a double tip (LDPE) packed in a cardboard container.
Pack size: 5 x 10 g with 5 cleaning towels. White intramammary syringe (LDPE) with a double tip (LDPE) packed in a cardboard container.
Pack size: 3 x 10 g with 3 cleaning towels. White intramammary syringe (LDPE) with a double tip (LDPE) packed in a cardboard container.

## Additional information

**Entitlement type:**

Marketing Authorisation

**Legal basis of product authorisation:**
Generic application (Article 13(1) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Vetcare Oy

**Marketing authorisation date:**
20/01/2021

**Manufacturing sites for batch release:**
KELA Kempisch Laboratorium Kela Laboratoria

**Responsible authority:**
Norwegian Medical Products Agency

**Authorisation number:**
19-13237

**Date of authorisation status change:**
21/01/2021

**Reference member state:**
Finland

**Procedure number:**
FI/V/0110/001

**Concerned member states:**
Austria Belgium Denmark Estonia France Germany Iceland Ireland Italy Latvia Lithuania Netherlands Norway Poland Spain Sweden United Kingdom (Northern Ireland)

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

# Documents

Summary of Product Characteristics

English (PDF)
Published on: 24/09/2025
Updated on: 13/03/2026
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Package Leaflet

English (PDF)
Published on: 16/01/2026
Updated on: 14/03/2026
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