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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:

Denmark

Available in:

Denmark

Package description:

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: 100 x 10 g with 100 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.
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.

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:
4/05/2021

Manufacturing sites for batch release:
KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:
Danish Medicines Agency

Authorisation number:
63786

Date of authorisation status change:
4/05/2021

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
Download

Package Leaflet

Labelling