

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:

Netherlands

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:

20/05/2016

Manufacturing sites for batch release:

KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 114527

Date of authorisation status change:

17/01/2022

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

Package Leaflet

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.

Labelling