PRURIVET N, Kožní roztok

Authorised

- Chloramphenicol
- Dexamethasone
- Benzyl benzoate

Product identification

Medicine name:
PRURIVET N, Kožní roztok

Active substance:
Chloramphenicol

Dexamethasone

Benzyl benzoate

Target species:
Dog

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Chloramphenicol
12.00 milligram(s) / 1.00 gram(s)

Dexamethasone

0.50 milligram(s) / 1.00 gram(s)

Benzyl benzoate
100.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Cutaneous solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD07CB04

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Czechia

Available in:
Czechia

Package description:
Available only in Czech
Available only in Czech

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetoquinol s.r.o.

Marketing authorisation date:
3/12/2003

Manufacturing sites for batch release:
Vetoquinol Biowet Sp. z o.o.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/069/03-C

Date of authorisation status change:
1/04/2015

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

Documents

Summary of Product Characteristics

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Package Leaflet

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

Labelling

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