NEOSTOMOSAN, Koncentrát pro kožní roztok

Authorised

- Tetramethrin
- Cypermethrin

Product identification

Medicine name:
NEOSTOMOSAN, Koncentrát pro kožní roztok

Active substance:
Tetramethrin

Cypermethrin

Target species:
Dog

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Tetramethrin
5.00 milligram(s) / 1.00 millilitre(s)

Cypermethrin
50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Concentrate for cutaneous solution

Withdrawal period by route of administration:

Cutaneous use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53AC30

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

Authorisation status:
Valid

Authorised in:
Czechia

Package description:
Available only in Czech
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:
Ceva Animal Health Slovakia s.r.o.

Marketing authorisation date:
8/07/1992

Manufacturing sites for batch release:

Ceva-Phylaxia Veterinary Biologicals Co. Ltd.
Chemark Kft.

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

Authorisation number:
96/444/92–C

Date of authorisation status change:
8/07/1992

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

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