

CORTIZEME, Kožní suspenze

Authorised

- NEOMYCIN SULFATE
- Prednisolone

Product identification

Medicine name:

CORTIZEME, Kožní suspenze

Active substance:

NEOMYCIN SULFATE

Prednisolone

Target species:

Cat

Dog

Route of administration:

Cutaneous use

Product details

Active substance and strength:

NEOMYCIN SULFATE

5000.00 international unit(s) / 1.00 millilitre(s)

Prednisolone

1.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Cutaneous suspension

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QD07CA03

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

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:

Virbac

Marketing authorisation date:

13/05/2003

Manufacturing sites for batch release:

Virbac

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/039/03-C

Date of authorisation status change:

27/07/2010

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

Documents

Summary of Product Characteristics

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

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.