File downloaded on 2026-11-09
Source URL: https://medicines.health.europa.eu/veterinary/en/600000078657

Cortizeme külsőleges szuszpenzió

Authorised

- Prednisolone
- NEOMYCIN SULFATE

Product identification

Medicine name:
Cortizeme külsőleges szuszpenzió

Active substance:
Prednisolone
NEOMYCIN SULFATE

Target species:
Dog
Cat

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Prednisolone
0.01 gram(s) / 100.00 millilitre(s)
NEOMYCIN SULFATE
5000.00 international unit(s) / 1.00 litre(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:
Hungary

Available in:
Hungary

Package description:
Available only in Hungarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:
Virbac

Marketing authorisation date:
13/11/2002

Manufacturing sites for batch release:
Virbac

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
13/11/2002

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.