Cortizeme, odos suspensija šunims ir katėms

Authorised

- NEOMYCIN SULFATE
- Prednisolone

Product identification

Medicine name:
Cortizeme, odos suspensija šunims ir katėms

Active substance:
NEOMYCIN SULFATE
Prednisolone

Target species:
Dog
Cat

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:
Lithuania

Available in:
Lithuania

Package description:
Available only in Lithuanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Virbac

Marketing authorisation date:
13/03/2001

Manufacturing sites for batch release:
Virbac

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/01/1234/001

Date of authorisation status change:
31/03/2011

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

Documents

Summary of Product Characteristics

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