

Cortizeme, nahasuspensioon koertele ja kassidele

Authorised

- NEOMYCIN SULFATE
- Prednisolone

Product identification

Medicine name:

Cortizeme, nahasuspensioon koertele ja kassidele

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:

Estonia

Available in:

Estonia

Package description:

Available only in Estonian

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:

12/04/2007

Manufacturing sites for batch release:

Virbac

Responsible authority:

State Agency Of Medicines

Authorisation number:

1466

Date of authorisation status change:

12/04/2007

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

Documents

Summary of Product Characteristics

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