Cortizeme 5000 IU/0,1 g cutaneous suspension

Authorised

- Prednisolone
- NEOMYCIN SULFATE

Product identification

Medicine name:
Кортизем 5000 IU/0,1 g суспензия за кожа
Cortizeme 5000 IU/0,1 g cutaneous suspension

Active substance:
Prednisolone
NEOMYCIN SULFATE

Target species:
Dog
Cat

Route of administration:
External use

Product details

Active substance and strength:
Prednisolone
0.10 gram(s) / 100.00 millilitre(s)

NEOMYCIN SULFATE
5000.00 international unit(s) / 100.00 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:
Bulgaria

Package description:
Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Virbac

Marketing authorisation date:
28/11/2010

Manufacturing sites for batch release:
Virbac

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1483

Date of authorisation status change:
28/11/2010

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 and Labelling

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