CORTIZEME EMULSION POUR APPLICATION CUTANEE

Authorised

- NEOMYCIN SULFATE
- Prednisolone

Product identification

Medicine name:
CORTIZEME EMULSION POUR APPLICATION CUTANEE

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 emulsion

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD07CA03

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
France

Available in:
France

Package description:
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Virbac

Marketing authorisation date:
2/02/1990

Manufacturing sites for batch release:
Virbac

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/6326271 1/1990

Date of authorisation status change:
2/02/2010

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

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