PREDNIDERM

Authorised

- Prednisolone acetate
- NEOMYCIN SULFATE

Product identification

Medicine name:
PREDNIDERM

Active substance:
Prednisolone acetate

NEOMYCIN SULFATE

Target species:
Dog
Cat
Horse

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Prednisolone acetate
1.34 milligram(s) / 1.00 gram(s)

NEOMYCIN SULFATE

333.33 international unit(s) / 1.00 gram(s)

Pharmaceutical form:
Cutaneous emulsion

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD07CA01

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

Authorisation status:
Valid

Authorised in:
Luxembourg

Available in:
Luxembourg

Package description:
Available only in French
Available only in French
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:
Vetoquinol S.A.

Marketing authorisation date:
19/09/1992

Manufacturing sites for batch release:

Vetoquinol S.A.

Responsible authority:
Ministry Of Health And Social Security

Authorisation number:
V 808/09/07/1232

Date of authorisation status change:
25/05/2012

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

Documents

Summary of Product Characteristics

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