DUPHADERM

Not authorised

- Hexetidine
- Prednisolone acetate

Product identification

Medicine name:
DUPHADERM

Active substance:
Hexetidine

Prednisolone acetate

Target species:
Dog
Cat

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Hexetidine
1.50 milligram(s) / 1.00 millilitre(s)

Prednisolone acetate
1.39 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Cutaneous solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD07BA01

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

Authorisation status:
Surrendered

Authorised in:
France

Package description:
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
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:

Zoetis France

Marketing authorisation date:
17/02/1992

Manufacturing sites for batch release:
Zoetis Manufacturing & Research Spain S.L.

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

Authorisation number:
FR/V/7689822 4/1992

Date of authorisation status change:
15/10/2025

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

Documents

Summary of Product Characteristics

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