Fungiderm 0,005 g/ml roztwór dla koni, lisów, psów, kotów, świnek morskich, myszy, szczurów i królików

Authorised

- Clotrimazole

Product identification

Medicine name:
Fungiderm 0,005 g/ml roztwór dla koni, lisów, psów, kotów, świnek morskich, myszy, szczurów i królików

Active substance:
Clotrimazole

Target species:
Rat
Mouse
Dog
Rabbit
Fox
Cat
Horse
Guinea pig

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Clotrimazole
0.01 gram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Cutaneous solution

Withdrawal period by route of administration:
Cutaneous use:

- **Rat**
  - All relevant tissues. no withdrawal period — Not applicable.
- **Mouse**
  - All relevant tissues. no withdrawal period — Not applicable.
- **Dog**
- **Rabbit**
  - Not applicable. 0 day
- **Fox**
  - All relevant tissues. no withdrawal period — Not applicable.
- **Cat**
- **Horse**

- Not applicable. 0 day

Do not use in horses whose tissues are intended for human consumption. Horses treated with the product must be identified as not intended for human consumption in the animal treatment book and on the identification document (passport) accompanying registered equines.

- **Guinea pig**
  - All relevant tissues. no withdrawal period — Not applicable.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD01AC01

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

Authorisation status:
Valid

Authorised in:
Poland

Package description:
Available only in Polish
Available only in Polish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Biowet Drwalew Sp. z o.o.

Marketing authorisation date:
21/05/1999

Manufacturing sites for batch release:
Drwalewskie Zaklady Przemyslu Bioweterynaryjnego S.A.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
0767

Date of authorisation status change:
21/05/1999

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

Documents

Package Leaflet

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

Summary of Product Characteristics

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

Labelling

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