

Fungidermin 20 mg/g Maść

Authorised

- Clotrimazole

Product identification

Medicine name:

Fungidermin 20 mg/g Maść

Active substance:

Clotrimazole

Target species:

Horse

Cat

Dog

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Clotrimazole

20.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Ointment

Withdrawal period by route of administration:**Cutaneous use:**

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Horse

- All relevant tissues. no withdrawal period

Do not use in horses whose tissues are intended for human consumption. Horses ever treated with Fungidermin must not be used for slaughter for human consumption. Only use in horses that have a signed declaration in their passport stating “not intended for slaughter for human consumption in accordance with current legislation”.

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

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:

30/06/2000

Manufacturing sites for batch release:

Drwalewskie Zakłady Przemysłu Bioweterynaryjnego S.A.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

1031

Date of authorisation status change:

30/06/2000

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

Documents

Package Leaflet

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Labelling

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.