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Insol Dermatophyton Zawiesina do wstrzykiwań

Authorised

- Trichophyton verrucosum, strain 410, Inactivated
- Trichophyton mentagrophytes, strain 551, Inactivated
- Trichophyton equinum, strain 381, Inactivated
- Trichophyton mentagrophytes, strain 1032, Inactivated
- Microsporum canis, strain 1393, Inactivated
- Microsporum distortum, strain 120, Inactivated
- Microsporum canis, strain 1311, Inactivated
- Nannizzia gypsea, strain 59, Inactivated

Product identification

Medicine name:
Insol Dermatophyton Zawiesina do wstrzykiwań

Active substance:
Trichophyton verrucosum, strain 410, Inactivated
Trichophyton mentagrophytes, strain 551, Inactivated
Trichophyton equinum, strain 381, Inactivated
Trichophyton mentagrophytes, strain 1032, Inactivated
Microsporum canis, strain 1393, Inactivated
Microsporum distortum, strain 120, Inactivated
Microsporum canis, strain 1311, Inactivated
Nannizzia gypsea, strain 59, Inactivated

Target species:
Cat
Horse
Dog

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Trichophyton verrucosum, strain 410, Inactivated
Trichophyton mentagrophytes, strain 551, Inactivated
Trichophyton equinum, strain 381, Inactivated
Trichophyton mentagrophytes, strain 1032, Inactivated
Microsporum canis, strain 1393, Inactivated
Microsporum distortum, strain 120, Inactivated
Microsporum canis, strain 1311, Inactivated
Nannizzia gypsea, strain 59, Inactivated

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Horse**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI05AQ02
QI06AQ01
QI07AQ01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Poland

Available in:
Poland

Package description:
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
29/04/2004

Manufacturing sites for batch release:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

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

Authorisation number:
1405

Date of authorisation status change:
29/04/2004

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

Documents

Labelling

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Summary of Product Characteristics

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

Package Leaflet

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