

Imaverol, 100mg/ml, Koncentrát pro kožní emulzi

Authorised

- Enilconazole

Product identification

Medicine name:

Imaverol, 100mg/ml, Koncentrát pro kožní emulzi

Active substance:

Enilconazole

Target species:

Horse

Dog

Cattle

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Enilconazole

100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Cutaneous emulsion

Withdrawal period by route of administration:

Cutaneous use:

•

Horse

- Meat and offal. 0 day
- Milk. 0 hour

•

Cattle

- Meat and offal. 0 day
 - Milk. 0 hour
-

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QD01AC90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Czechia

Package description:

Available only in [Czech](#)

Available only in [Czech](#)

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:

Audevard

Marketing authorisation date:

9/09/1992

Manufacturing sites for batch release:

Lusomedicamenta Sociedade Tecnica Farmaceutica S.A.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/519/92-S/C

Date of authorisation status change:

17/04/2009

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.

Package Leaflet

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.