

Imaverol Vet. koncentrat til kutanopløsning 100 mg/ml

Authorised

- Enilconazole
- Enilconazole

Product identification

Medicine name:

Imaverol Vet. koncentrat til kutanopløsning 100 mg/ml

Active substance:

Enilconazole

Enilconazole

Target species:

Cattle

Dog

Horse

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Enilconazole

100.00 milligram(s) / 1.00 millilitre(s)

Enilconazole
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Concentrate for cutaneous solution

Withdrawal period by route of administration:

Cutaneous use:

-

Cattle

- Meat and offal. 3 day
- Meat and offal. 3 day
- Milk. 1 day
- Milk. 1 day

-

Horse

- Meat and offal. 3 day
 - Meat and offal. 3 day
-

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QD01AC90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

Package description:

Available only in Danish

Available only in Danish

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Audevard

Marketing authorisation date:

21/10/1983

Manufacturing sites for batch release:

Lusomedicamenta Sociedade Tecnica Farmaceutica S.A.

Janssen Cilag Farmaceutica Lda.

Responsible authority:

Danish Medicines Agency

Authorisation number:

10562

Date of authorisation status change:

21/10/1983

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.