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
- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day

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:
9/09/1992

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

Documents

Summary of Product Characteristics

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Package Leaflet

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Labelling

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