IMAVEROL

Authorised

- Enilconazole

Product identification

Medicine name:
ИМАВЕРОЛ
IMAVEROL

Active substance:
Enilconazole

Target species:
Cattle
Horse
Dog

Route of administration:
Pour-on use

Product details

Active substance and strength:
Enilconazole
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Cutaneous solution

Withdrawal period by route of administration:

Pour-on use:

- **Cattle**
  - Meat and offal. no withdrawal period

Говеда и коне: Месо и вътрешни органи: нула дни. Мляко: нула дни

- **Horse**
- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD01AC90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian
Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Audevard

Marketing authorisation date:
28/10/2008

Manufacturing sites for batch release:
Lusomedicamenta Sociedade Tecnica Farmaceutica S.A.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2107

Date of authorisation status change:
13/10/2019

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 and Labelling

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

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