Imaverol 100 mg/mL, δερματικό διάλυμα, πυκνό διάλυμα για αραίωση για βοοειδή, άλογα, σκύλους

Authorised

- Enilconazole

Product identification

Medicine name:
Imaverol 100 mg/mL, δερματικό διάλυμα, πυκνό διάλυμα για αραίωση για βοοειδή, άλογα, σκύλους

Active substance:
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)

Pharmaceutical form:
Concentrate for cutaneous solution

Withdrawal period by route of administration:
Cutaneous use:

- **Cattle**
  - Meat and offal. 0 day
- **Dog**
  - Not applicable. no withdrawal period
- **Horse**
  - Meat and offal. 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:
Greece

Available in:
Greece

Package description:
Available only in Greek
Available only in Greek

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:
23/07/2018

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

Responsible authority:
National Organization For Medicines

Authorisation number:
82146/29-07-2022/09-02-2024/K-0234301

Date of authorisation status change:
8/02/2024

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