RAFOXANIDE/PROVET 300MG/TAB ΔΙΣΚΙΟ

Awtorizzat

- Rafoxanide

Product identification

Isem tal-mediċina:
RAFOXANIDE/PROVET 300MG/TAB ΔΙΣΚΙΟ

Sustanza attiva:
Disponibbli biss fi English

Speċi li fuqhom ser jintuża l-prodott:
Disponibbli biss fi Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Swedish Icelandic Norwegian

Metodu ta' amministrazzjoni:
Użu orali

Product details

Sustanza attiva / Qawwa:
Disponibbli biss fi English
300.00 milligram(s) / 1.00 Tablet

Forma farmaċewtika:
Pillola

Withdrawal period by route of administration:

Użu orali:

- **Sheep**

  - Meat and offal. 42 day

ΓΑΛΑ: Δεν επιτρέπεται η χρήση σε θηλθκά πρόβατα που παράγουν γάλα για ανθρώπινη κατανάλωση, περιλαμβανομένης της ξηράς περιόδου και να μην χρησιμοποιείται για ένα έτος πριν από τον πρώτο τοκετό

Kodiċi veterinarju anatomiku terapewtiku kimiku (ATCvet):
QP52AG05

Status legali tal-provvista:
Disponibbli biss fi Czech Estonian English French Italian Latvian Portuguese Slovenian Finnish Swedish Icelandic Norwegian

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Status tal-awtorizzazzjoni:
Valid

Authorised in:
Disponibbli biss fi Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Deskrizzjoni tal-pakkett:
Disponibbli biss fi Greek

Additional information

Entitlement type:
Disponibbli biss fi English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

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Bażi ġuridika tal-awtorizzazzjoni tal-prodott:
Disponibbli biss fi English

Id-detentur tal-awtorizzazzjoni għat-tqegħid fis-suq:
Provet S.A.

Marketing authorisation date:

28/03/1989

Siti ta’ manifattura b’rilaxx tal-lott:
Provet S.A.

Awtorità responsabbli:
National Organization For Medicines

Numru tal-awtorizzazzjoni:
89517/15-12-2009/K-0041601

Data tal-bidla fl-istatus tal-awtorizzazzjoni:
11/10/2021

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

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