Noromectin Drench 0.8 mg/ml Oral Solution for Sheep

Authorised

- Ivermectin

Product identification

Medicine name:
NOROMECTIN DRENCH 0.8MG/ML ΠΟΣΙΜΟ ΔΙΑΛΥΜΑ ΓΙΑ ΠΡΟΒΑΤΑ
Noromectin Drench 0.8 mg/ml Oral Solution for Sheep

Active substance:
Ivermectin

Target species:
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Ivermectin
0.80 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Withdrawal period by route of administration:

Oral use:

- **Sheep**
  - Meat and offal. 10 day
  - Milk. no withdrawal period

NOT AUTHORIZED FOR USE IN LACTATING ANIMALS PRODUCING MILK FOR HUMAN CONSUPMTION

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA01

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

Authorisation status:
Valid

Authorised in:
Cyprus

Package description:
Available only in Greek
Available only in Greek

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:
Spyros Stavrinides Chemicals Limited

Marketing authorisation date:

30/10/2000

Manufacturing sites for batch release:
Norbrook Laboratories Limited

Responsible authority:
Ministry Of Agriculture Rural Development And Environment

Authorisation number:
19056

Date of authorisation status change:
31/05/2012

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.