VERMAX D

Authorised

- Ivermectin
- Closantel sodium dihydrate

Product identification

Medicine name:
VERMAX D

Active substance:
Ivermectin

Closantel sodium dihydrate

Target species:
Sheep

Route of administration:
Subcutaneous use

Product details

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

Closantel sodium dihydrate
135.90 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Sheep**
  - Meat and offal. 28 day
  - Milk. no withdrawal period No withdrawal period

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA51

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription except for some pack sizes

Authorisation status:
Valid

Authorised in:
France

Package description:
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
26/03/2010

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/9829328 1/2010

Date of authorisation status change:
19/05/2015

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.