MOLEMEC PLUS PASTE

Authorised

- Ivermectin
- Praziquantel

Product identification

Medicine name:
MOLEMEC PLUS PASTE

Active substance:
Ivermectin

Praziquantel

Target species:
Horse

Route of administration:
Oral use

Product details

Active substance and strength:
Ivermectin
15.50 milligram(s) / 1.00 gram(s)

Praziquantel
77.50 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Oral paste

Withdrawal period by route of administration:

Oral use:

- **Horse**

  - Milk. no withdrawal period

Do not use in mares producing milk for human consumption.

  - Meat and offal. 30 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA51

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

Box of 1 sachet of 1 syringe of 7,74 g
Box of 50 sachets of 1 syringe of 9,68 g
Box of 50 sachets of 1 syringe of 7,74 g
Box of 1 sachet of 1 syringe of 14,19 g
Box of 1 sachet of 1 syringe of 9,68 g
Box of 50 sachets of 1 syringe of 14,19 g

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:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
9/07/2012

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France SCS

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

Authorisation number:
FR/V/0334369 6/2012

Date of authorisation status change:
12/09/2016

Reference member state:
France

Procedure number:
FR/V/0361/001

Concerned member states:
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

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eu-puar-frv0361001-mr-rpe409-en.pdf