

FUREXEL COMBI ORAL PASTE

Authorised

- Ivermectin
- Praziquantel

Product identification

Medicine name:

FUREXEL COMBI ORAL PASTE

Furexel Combi Pasta voor oraal gebruik

Furexel Combi Pâte orale

Furexel Combi Paste zum Einnehmen

Active substance:

Ivermectin

Praziquantel

Target species:

Horse

Route of administration:

Oral use

Product details

Active substance and strength:

Ivermectin

15.50 gram(s) / 1.00 gram(s)

Praziquantel

77.50 gram(s) / 1.00 gram(s)

Pharmaceutical form:

Oral paste

Withdrawal period by route of administration:**Oral use:**

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Horse

- Meat and offal. 30 day
- Milk. no withdrawal period

Do not use in mares producing milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA51

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Available in:

Belgium

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:

Boehringer Ingelheim Animal Health Belgium S.A.

Marketing authorisation date:

6/02/2006

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V279221

Date of authorisation status change:

6/02/2006

Reference member state:

France

Procedure number:

FR/V/0360/001

Concerned member states:

Belgium Germany Luxembourg United Kingdom (Northern Ireland)

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.

Package Leaflet

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Labelling

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