# Milteforan 20 mg/ml oral solution for dogs

Authorised

Miltefosine

# Product identification

#### **Medicine name:**

Milteforan 20 mg/ml orale oplossing voor honden Milteforan 20 mg/ml oral solution for dogs

#### **Active substance:**

Miltefosine

## **Target species:**

Dog

### **Route of administration:**

Oral use

# **Product details**

## **Active substance and strength:**

Miltefosine

20.00 milligram(s) / 1.00 millilitre(s)

#### **Pharmaceutical form:**

Oral solution

## Withdrawal period by route of administration:

#### Oral use:

•

Dog

## **Anatomical therapeutic chemical veterinary (ATCvet) codes:**

**QP51DX07** 

## Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### Authorised in:

**Netherlands** 

## Package description:

Carton box containing a 90 ml Polyethylene terephthalate (PET) bottle equipped with a sampling polypropylene (PP) snap cap with silicone stopper and a 3 ml polypropylene (PP) syringe graduated every 0.1 ml

Carton box containing a 60 ml Polyethylene terephthalate (PET) bottle equipped with a sampling polypropylene (PP) snap cap with silicone stopper and a 3 ml polypropylene (PP) syringe graduated every 0.1 ml

Carton box containing a 30 ml Polyethylene terephthalate (PET) bottle equipped with a sampling polypropylene (PP) snap cap with silicone stopper and a 3 ml polypropylene (PP) syringe graduated every 0.1 ml

# Additional information

# **Entitlement type:**

Marketing Authorisation

## Legal basis of product authorisation:

Generic application (Article 18 of Regulation (EU) 2019/6)

# Marketing authorisation holder:

Virbac

Marketing authorisation date: 21/02/2024
Manufacturing sites for batch release: Virbac
Responsible authority: Medicines Evaluation Board
Authorisation number: REG NL 130834
Date of authorisation status change: 26/02/2024
Reference member state: Netherlands
Procedure number: NL/V/0405/001
<b>Concerned member states:</b> Bulgaria Croatia Cyprus France Greece Italy Portugal Romania Slovenia Spain
Generic of: 60000004401
To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>
Documents
Combined File of all Documents
English (PDF)

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