Source URL: https://medicines.health.europa.eu/veterinary/en/600000993033

Milteforan 20 mg/ml oral solution for dogs

Authorised

Miltefosine

Product identification

Medicine name:

Milteforan 20 mg/ml oral solution for dogs Milteforan 20 mg/ml πόσιμο διάλυμα για σκύλους

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:

Cyprus

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

11/03/2024
Manufacturing sites for batch release: Virbac
Responsible authority: Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment
Authorisation number: CY00938V
Date of authorisation status change: 11/03/2024
Reference member state: Netherlands
Procedure number: NL/V/0405/001
Concerned member states: Bulgaria Croatia Cyprus France Greece Italy Portugal Romania Slovenia Spain
Generic of: 60000004401
To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

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