

# Eradia 125 mg/ml, oral suspension for dogs

Authorised

- Metronidazole

## Product identification

**Medicine name:**

Eradia 125 mg/ml, oral suspension for dogs

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**Active substance:**

Metronidazole

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**Target species:**

Dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Metronidazole

125.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Oral suspension

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

QJ01XD01

QP51AA01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Cyprus

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**Available in:**

Cyprus

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**Package description:**

30 ml presentation: white opaque polyethylene terephthalate (PET) bottle equipped with a sampling polypropylene (PP) snap cap with silicon stopper and a 3 ml polypropylene (PP) syringe placed in a carton box

100 ml presentation: white opaque polyethylene terephthalates (PET) bottle equipped with a sampling Polyethylene (PE) screw cap with PE seal and a 3 ml polypropylene (PP) oral syringe placed in a carton box

30 ml presentation: white opaque polyethylene terephthalate (PET) bottle equipped with a sampling polyethylene (PE) screw cap with PE seal and a 3 ml polypropylene (PP) oral syringe placed in a carton box

100 ml presentation: white opaque polyethylene terephthalate (PET) bottle equipped with a sampling polypropylene (PP) snap cap with silicon stopper and a 3 ml polypropylene (PP) syringe placed in a carton box

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Virbac

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**Marketing authorisation date:**

25/09/2018

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**Manufacturing sites for batch release:**

Delpharm Huningue S.A.S.

Virbac

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**Responsible authority:**

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

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**Authorisation number:**

CY00703V

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**Date of authorisation status change:**

25/09/2018

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0232/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania  
Luxembourg Poland Portugal Romania Slovakia Slovenia Spain Sweden  
United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
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## Documents

Combined File of all Documents

English (PDF)

Published on: 9/05/2024

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