

# AFILARIA SR 3.4 mg/ml powder and solvent for suspension for injection for dogs

Authorised

- Moxidectin

## Product identification

**Medicine name:**

AFILARIA SR 3.4 mg/ml powder and solvent for suspension for injection for dogs

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**Active substance:**

Moxidectin

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**Target species:**

Dog

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Moxidectin

100.00 milligram(s)/gram / 1.00 gram(s)

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**Pharmaceutical form:**

Powder and solvent for suspension for injection

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

QP54AB02

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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:**

Portugal

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**Available in:**

Portugal

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**Package description:**

Cardboard box with 1 powder vial containing 197,3 mg, 1 solvent vial with 5,67 ml and 1 adapter for reconstitution and withdrawal, 1 syringe and 1 needle.

Cardboard box with 1 powder vial containing 197,3 mg, 1 solvent vial with 5,67 ml and 1 adapter for reconstitution and withdrawal.

Cardboard box with 1 powder vial containing 592 mg, 1 solvent vial with 17 ml and 1 adapter for reconstitution and withdrawal, 1 syringe and 1 needle.

Cardboard box with 1 powder vial containing 592 mg, 1 solvent vial with 17 ml and 1 adapter for reconstitution and withdrawal.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Fatro S.p.A.

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**Marketing authorisation date:**

5/07/2019

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**Manufacturing sites for batch release:**

Fatro S.p.A.

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**Responsible authority:**

Directorate General For Food And Veterinary

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**Authorisation number:**

1277/01/19DFVPT

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**Date of authorisation status change:**

20/06/2024

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0315/001

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**Concerned member states:**

Austria Bulgaria Croatia Czechia France Greece Hungary Italy Portugal  
Romania Slovakia Slovenia

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

Package Leaflet

Labelling

Combined File of all Documents

English (PDF)

Published on: 3/12/2025

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