

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

Active substance:

Moxidectin

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Moxidectin

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

Pharmaceutical form:

Powder and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AB02

Legal status of supply:

This information is not available for this product.

Authorisation status:

Valid

Authorised in:

Czechia

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.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Fatro S.p.A.

Marketing authorisation date:

8/06/2022

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/020/22-C

Date of authorisation status change:

8/06/2022

Reference member state:

Spain

Procedure number:

ES/V/0315/001

Concerned member states:

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

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