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
AFILARIA SR 3,4 mg/ml por és oldószer szuszpenziós injekcióhoz kutyák számára
A.U.V.

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

Withdrawal period by route of administration:

Subcutaneous use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Hungary

Available in:

Hungary

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:
7/12/2021

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
4297/X/21 NÉBIH ÁTI (1 db 592 mg port tartalmazó ü

Date of authorisation status change:
7/12/2021

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

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)

Published on: 12/04/2023
Download

Package Leaflet

Labelling

eu-PUAR-afilaria-sr-3.4-mg-ml-powder-and-solvent-for-suspension-for-injection-for-dogs-en.pdf

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