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, prašak i otapalo za suspenziju za injekciju, za pse

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:

Croatia

Available in:

Croatia

Package description:

Available only in Croatian

Available only in Croatian

Available only in Croatian

Available only in Croatian

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:

14/03/2022

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

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/22-01/149

Date of authorisation status change:
26/08/2022

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

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

Documents

Summary of Product Characteristics

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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