

Alizin 30 mg/ml Solution for Injection

Authorised

- Aglepristone

Product identification

Medicine name:

Alizin 30 mg/ml Solution for Injection
Alizin 30 mg/ml Oplossing voor injectie
Alizin 30 mg/ml Solution injectable
Alizin 30 mg/ml Injektionslösung

Active substance:

Aglepristone

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Aglepristone
30.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG03XB90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Available in:

Belgium

Package description:

Vial (glass, type II) of 5 ml for injectable preparation, with bromobutyl stopper and aluminum cap.Presentation:- box of 1 vial of 5ml

Vial (glass, type II) of 10 ml for injectable preparation, with bromobutyl stopper and aluminum cap.Presentation:- box of 1 vial of 10ml.

Vial (glass, type II) of 30 ml for injectable preparation, with bromobutyl stopper and aluminum cap.Presentation:- box of 1 vial of 30ml.

Vials (glass, type II) of 10 ml for injectable preparations, with bromobutyl stoppers and aluminum caps.Presentation:- box of 10 vials of 10ml.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Virbac

Marketing authorisation date:

14/06/2004

Manufacturing sites for batch release:

Virbac

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V263986

Date of authorisation status change:

14/06/2004

Reference member state:

Ireland

Procedure number:

IE/V/0500/001

Concerned member states:

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

To consult adverse reactions on veterinary medicinal products please go to

www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)

Published on: 16/11/2025

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Package Leaflet

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Labelling

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Combined File of all Documents