

# Alizin 30 mg/ml Solution for Injection

Authorised

- Aglepristone

## Product identification

**Medicine name:**

Alizin 30 mg/ml Solution for Injection

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**Active substance:**

Aglepristone

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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:**

Aglepristone

30.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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

QG03XB90

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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:**

Bulgaria

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**Available in:**

Bulgaria

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**Package description:**

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

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 5 ml for injectable preparation, with bromobutyl stopper and aluminum cap.Presentation:- box of 1 vial of 5ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Virbac

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**Marketing authorisation date:**

29/06/2011

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**Manufacturing sites for batch release:**

Virbac

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**Responsible authority:**

Bulgarian Food Safety Authority

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**Authorisation number:**

0022-1604

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**Date of authorisation status change:**

29/06/2011

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0500/001

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**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)

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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

English (PDF)

Published on: 16/11/2025

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## Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.

## Combined File of all Documents

## Package Leaflet

This document does not exist in this language (English). You can find it in another language below.