

ACEGON 50 mcg/ml solution for injection

Authorised

- Gonadorelin acetate

Product identification

Medicine name:

ACEGON 50 mcg/ml solution for injection

Active substance:

Gonadorelin acetate

Target species:

Cattle (cow)

Cattle (heifer)

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Gonadorelin acetate

50.00 microgram(s)/millilitre / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH01CA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Available in:

Netherlands

Package description:

cardboard box containing 1 vial of 6 ml

cardboard box containing 1 vial of 20 ml

cardboard box containing 1 vial of 50 ml

cardboard box containing 1 vial of 100 ml

cardboard box containing 10 vials of 6 ml

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:

Laboratorios Syva S.A.

Marketing authorisation date:

14/02/2011

Manufacturing sites for batch release:

Laboratorios Syva S.A.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 107265

Date of authorisation status change:

17/01/2022

Reference member state:

Spain

Procedure number:

ES/V/0158/001

Concerned member states:

Austria Belgium Cyprus Czechia France Greece Hungary Ireland Italy
Luxembourg Netherlands Poland Portugal Slovakia
United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)

Published on: 24/07/2025

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Summary of Product Characteristics

Package Leaflet

Labelling

eu-PUAR-acegon-50-mcg-ml--solution-for-injection-en.pdf