File downloaded on 2026-11-03
Source URL: https://medicines.health.europa.eu/veterinary/en/600000016882

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:
Ireland

Available in:
Ireland

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:
28/02/2014

Manufacturing sites for batch release:
Laboratorios Syva S.A.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10495/004/001

Date of authorisation status change:
28/02/2014

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

Summary of Product Characteristics

Package Leaflet

Labelling

Combined File of all Documents

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