ARVILAP

Suspended

- Rabbit haemorrhagic disease virus, type 1, strain LO1, Inactivated

Product identification

Medicine name:
ARVILAP

Active substance:
Rabbit haemorrhagic disease virus, type 1, strain LO1, Inactivated

Target species:
Rabbit

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Rabbit haemorrhagic disease virus, type 1, strain LO1, Inactivated
80.00 haemagglutination inhibiting unit(s) / 1.00 Dose

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Rabbit**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI08AA01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Suspended

Authorised in:
Spain

Package description:
Available only in Spanish
Available only in Spanish
Available only in Spanish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Laboratorios Ovejero S.A.U.

Marketing authorisation date:
19/04/1989

Manufacturing sites for batch release:
Laboratorios Ovejero S.A.U.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
2947 ESP

Date of authorisation status change:
1/01/2023

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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