

Gumboriffa ενέσιμο γαλάκτωμα για όρνιθες

Authorised

- Infectious bursal disease virus, strain VNJO, Inactivated

Product identification

Medicine name:

Gumboriffa ενέσιμο γαλάκτωμα για όρνιθες

Active substance:

Infectious bursal disease virus, strain VNJO, Inactivated

Target species:

Poultry

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Infectious bursal disease virus, strain VNJO, Inactivated

1000000.00 50% cell culture infectious dose / 0.30 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:**Intramuscular use:**

-

Poultry

- Meat and offal. 0 day

Subcutaneous use:

-

Poultry

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Available in:

Greece

Package description:

Available only in Greek

Available only in Greek

Available only in Greek

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone)

Marketing authorisation holder:

Boehringer Ingelheim Animal Health France

Marketing authorisation date:

5/07/1990

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

National Organization For Medicines

Authorisation number:

18565/14-02-2024/K-0125801

Date of authorisation status change:

13/02/2024

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.