GALLIMUNE 403 ND+IB+IBD+REO, injekcinė emulsija

Not authorised

- Newcastle disease virus, strain Ulster 2C, Inactivated
- Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Infectious bursal disease virus, strain VNJO, Inactivated
- Avian reovirus, strain S1133, Inactivated

Product identification

Medicine name:
GALLIMUNE 403 ND+IB+IBD+REO, injekcinė emulsija

Active substance:
Newcastle disease virus, strain Ulster 2C, Inactivated
Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
Infectious bursal disease virus, strain VNJO, Inactivated
Avian reovirus, strain S1133, Inactivated

Target species:
Chicken

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:
Newcastle disease virus, strain Ulster 2C, Inactivated
100000000.00 50% Embryo Infective Dose / 1.00 Dose

Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
5011870.00 50% Embryo Infective Dose / 1.00 Dose

Infectious bursal disease virus, strain VNJO, Inactivated
501187.00 50% cell culture infectious dose / 1.00 Dose

Avian reovirus, strain S1133, Inactivated
10000000.00 50% cell culture infectious dose / 1.00 Dose

Pharmaceutical form:
Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AA16

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription except for some pack sizes

Authorisation status:
Surrendered

Authorised in:
Lithuania

Package description:
Available only in Lithuanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Boehringer Ingelheim Animal Health Italia S.p.A. In Breve Boehringer Ingelheim Ah It S.p.A.

Marketing authorisation date:
18/11/1999

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health Italia S.p.A. In Breve Boehringer Ingelheim Ah It S.p.A.

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/99/1006/001

Date of authorisation status change:
10/04/2010

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.