GALLIMUNE 407 ND+IB+EDS+ART

Authorised

- Newcastle disease virus, strain Ulster 2C, Inactivated
- Eggdrop syndrome-1976 virus, strain V127, Inactivated
- Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Turkey rhinotracheitis virus, strain VCO3, Inactivated

Product identification

Medicine name:
GALLIMUNE 407 ND+IB+EDS+ART

Active substance:
Newcastle disease virus, strain Ulster 2C, Inactivated

Eggdrop syndrome-1976 virus, strain V127, Inactivated

Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Turkey rhinotracheitis virus, strain VCO3, Inactivated

Target species:
Chicken (layer hen)
Chicken (for reproduction)

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Newcastle disease virus, strain Ulster 2C, Inactivated
10.00 haemagglutination inhibiting unit(s) / 1.00 Dose

Eggdrop syndrome-1976 virus, strain V127, Inactivated
162.00 haemagglutination inhibiting unit(s) / 1.00 Dose

Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
10.00 haemagglutination inhibiting unit(s) / 1.00 Dose

Turkey rhinotracheitis virus, strain VCO3, Inactivated
60.00 interference percentage unit(s) / 1.00 Dose

Pharmaceutical form:

Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA18

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Bulgaria

Package description:

Available only in Bulgarian

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

Marketing authorisation date:
12/03/2007

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

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2016

Date of authorisation status change:
14/04/2013

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.