

GALLIMUNE 407

Authorised

ND+IB+EDS+ART Water-in oil emulsion for injection

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

Product identification

Medicine name:

GALLIMUNE 407 ND+IB+EDS+ART Water-in oil emulsion for injection

Active substance:

Turkey rhinotracheitis virus, strain VCO3, Inactivated

Eggdrop syndrome-1976 virus, strain V127, Inactivated

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

Newcastle disease virus, strain Ulster 2C, Inactivated

Target species:

Chicken (for reproduction)

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Turkey rhinotracheitis virus, strain VCO3, Inactivated
0.76 interference percentage unit(s) / 0.30 millilitre(s)

Eggdrop syndrome-1976 virus, strain V127, Inactivated
180.00 log₁₀ haemagglutination inhibiting unit(s) / 0.30 millilitre(s)

Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
18.00 log₁₀ haemagglutination inhibiting unit(s) / 0.30 millilitre(s)

Newcastle disease virus, strain Ulster 2C, Inactivated
50.00 50% Protective Dose / 0.30 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Chicken (for reproduction)

- Egg. 0 day
 - Meat and offal. 0 day
-

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA18

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Germany

Package description:

(ID4): 1 Box with 10 Bottle (PolyPropylene) with 300 millilitre(s) (3000 millilitre(s))

(ID3): 1 Box with 1 Bottle (PolyPropylene) with 300 millilitre(s) (300 millilitre(s))
(ID2): 1 Box with 10 Bottle (PolyPropylene) with 150 millilitre(s) (1500 millilitre(s))
(ID1): 1 Box with 1 Bottle (PolyPropylene) with 150 millilitre(s) (150 millilitre(s))

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:

Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:

11/05/2000

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

Paul-Ehrlich-Institut

Authorisation number:

PEI.V.01043.01.1

Date of authorisation status change:

29/04/2009

Reference member state:

Germany

Procedure number:

DE/V/0229/001

Concerned member states:

Austria Belgium Cyprus Czechia Denmark Finland France Greece Hungary

Ireland Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal
Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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