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

Authorised

- Turkey rhinotracheitis virus, strain VCO3, Inactivated
- Eggdrop syndrome-1976 virus, strain V127, Inactivated
- 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

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)

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

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Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

Package description:
(ID4): 1 Box with 10 Bottle (PolyPropylene) with 300 millilitre(s) (3000 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))
(ID3): 1 Box with 1 Bottle (PolyPropylene) with 300 millilitre(s) (300 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 Animal Health Denmark A/S

Marketing authorisation date:
29/04/2005

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Danish Medicines Agency

Authorisation number:
36564

Date of authorisation status change:
29/04/2005

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

Summary of Product Characteristics

English (PDF)
Published on: 28/01/2022
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Combined File of all Documents