

GALLIMUNE 302 ND+IB+EDS

Water-in oil emulsion for injection

Not
authorised

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

Active substance:

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:

Eggdrop syndrome-1976 virus, strain V127, Inactivated

162.00 log₁₀ haemagglutination inhibiting unit(s) / 0.30 millilitre(s)

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

10.00 log₁₀ haemagglutination inhibiting unit(s) / 0.30 millilitre(s)

Newcastle disease virus, strain Ulster 2C, Inactivated

10.00 log₁₀ haemagglutination inhibiting unit(s) / 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:

QI01AA13

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Czechia

Package description:

(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))

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

Marketing authorisation date:

8/09/2004

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

97/137/04-C

Date of authorisation status change:

28/08/2024

Reference member state:

Germany

Procedure number:

DE/V/0227/001

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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Package Leaflet

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Labelling

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