

GALLIMUNE 303 ND+IB+ART

Water-in oil emulsion for injection

Not
authorised

- Turkey rhinotracheitis virus, strain VCO3, Inactivated
- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Newcastle disease virus, strain Ulster 2C, Inactivated

Product identification

Medicine name:

GALLIMUNE 303 ND+IB+ART Water-in oil emulsion for injection

Active substance:

Turkey rhinotracheitis virus, strain VCO3, 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)

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
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA21

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Italy

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

Marketing authorisation date:

21/05/2024

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

Ministry Of Health

Authorisation number:

103715

Date of authorisation status change:

28/07/2009

Reference member state:

Germany

Procedure number:

DE/V/0228/001

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

Documents

Summary of Product Characteristics

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.