Source URL: https://medicines.health.europa.eu/veterinary/en/600000027402

Gallimune 302 ND + IB + EDS, emulsie injectabilă

Not authorised

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

Product identification

Medicine name:

Gallimune 302 ND + IB + EDS, emulsie injectabilă

Active substance:

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

Target species:

Chicken

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Newcastle disease virus, strain Ulster 2C, Inactivated 10.00 haemagglutination inhibiting unit(s) / 0.30 millilitre(s)

Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated 10.00 haemagglutination inhibiting unit(s) / 0.30 millilitre(s)

Eggdrop syndrome-1976 virus, strain V127, Inactivated 162.00 haemagglutination inhibiting unit(s) / 0.30 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration: Intramuscular use:

Chicken

- Meat and offal. 0 day

The product is not administered during the laying period.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA13

Legal status of supply:

This information is not available for this product.

Authorisation status:

Surrendered

Authorised in:

Romania

Package description:

Available only in Romanian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone) - 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:

25/07/2006

Manufacturing sites for batch release:

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

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

110323

Date of authorisation status change:

31/07/2024

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.