GALLIMUNE 407 ND+IB+EDS+ART, emulzija za injekciju, za kokoši

Authorised

- 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, emulzija za injekciju, za kokoši

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 (layer hen)
Chicken (for reproduction)

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Turkey rhinotracheitis virus, strain VCO3, Inactivated
0.76 log10 optical density50 / 0.30 millilitre(s)

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

Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
18.00 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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA18

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Croatia

Available in:

Croatia

Package description:

Available only in Croatian
Available only in Croatian
Available only in Croatian
Available only in Croatian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
16/10/2018

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/15-01/371

Date of authorisation status change:
29/04/2024

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

Documents

Summary of Product Characteristics

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