ORNIVAC ND+IB2+EDS, injekční emulze pro kura domácího, Injekční emulze

Authorised

- Newcastle disease virus, strain SL-93, Inactivated
- Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Avian infectious bronchitis virus, strain D274, Inactivated
- Eggdrop syndrome-1976 virus, Inactivated

Product identification

Medicine name:
ORNIVAC ND+IB2+EDS, injekční emulze pro kura domácího, Injekční emulze

Active substance:
Newcastle disease virus, strain SL-93, Inactivated

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

Avian infectious bronchitis virus, strain D274, Inactivated

Eggdrop syndrome-1976 virus, Inactivated

Target species:
Chicken (pullet)

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Newcastle disease virus, strain SL-93, Inactivated
4.00 log2 haemagglutination inhibiting unit(s) / 1.00 Dose

Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
6.20 log2 haemagglutination inhibiting unit(s) / 1.00 Dose

Avian infectious bronchitis virus, strain D274, Inactivated
6.30 log2 haemagglutination inhibiting unit(s) / 1.00 Dose

Eggdrop syndrome-1976 virus, Inactivated
6.50 log2 haemagglutination inhibiting unit(s) / 1.00 Dose

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Chicken (pullet)**
  - Meat and offal. 0 day
  - Egg. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA13

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Czechia

Package description:

Available only in Czech

Available only in Czech
Available only in Czech
Available only in Czech

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:
Bioveta a.s.

Marketing authorisation date:
13/01/2014

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/003/14-C

Date of authorisation status change:
28/01/2022

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.

Package Leaflet

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

Labelling

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