Adenipravac-ND/IB Emulsja do wstrzykiwań

Authorised

- Newcastle disease virus, strain La Sota, Inactivated
- Infectious bronchitis virus, strain H52, Inactivated
- Eggdrop syndrome-1976 virus, strain V127, Inactivated

Product identification

Medicine name:
Adenipravac-ND/IB Emulsja do wstrzykiwań

Active substance:
Newcastle disease virus, strain La Sota, Inactivated

Infectious bronchitis virus, strain H52, Inactivated

Eggdrop syndrome-1976 virus, strain V127, Inactivated

Target species:
Chicken (hen)

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:

Newcastle disease virus, strain La Sota, Inactivated
100000000.00 50% Embryo Infective Dose / 0.50 millilitre(s)

Infectious bronchitis virus, strain H52, Inactivated
1000000.00 50% Embryo Infective Dose / 0.50 millilitre(s)

Eggdrop syndrome-1976 virus, strain V127, Inactivated
500.00 haemagglutinating units / 0.50 millilitre(s)

Pharmaceutical form:
Emulsion for injection/infusion

Withdrawal period by route of administration:

Subcutaneous use:

- **Chicken (hen)**
  - All relevant tissues. 0 day

Intramuscular use:

- **Chicken (hen)**
  - All relevant tissues. 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:
Poland

Package description:
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish

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:
Laboratorios Hipra S.A.

Marketing authorisation date:
22/11/2001

Manufacturing sites for batch release:
Laboratorios Hipra S.A.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
1241

Date of authorisation status change:
22/11/2001

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

Documents

Package Leaflet

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Labelling

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

Summary of Product Characteristics

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