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CEVAC ND IB EDS K

Not authorised

- Eggdrop syndrome-1976 virus, strain B8/78, Inactivated
- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Newcastle disease virus, strain La Sota, Inactivated

Product identification

Medicine name:
CEVAC ND IB EDS K

Active substance:
Eggdrop syndrome-1976 virus, strain B8/78, Inactivated

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Newcastle disease virus, strain La Sota, Inactivated

Target species:
Chicken
Chicken (hen)

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:

Eggdrop syndrome-1976 virus, strain B8/78, Inactivated
7.00 log2 haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
6.00 log2 haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Newcastle disease virus, strain La Sota, Inactivated
4.00 log2 haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Chicken**
  - Meat and offal. 0 day
  - Egg. 0 day

Intramuscular use:

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

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
Available only in Romanian
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:
Ceva Sante Animale Romania S.R.L.

Marketing authorisation date:
17/11/2009

Manufacturing sites for batch release:
Ceva-Phylaxia Zrt.

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
150006

Date of authorisation status change:
5/12/2023

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

Documents

Summary of Product Characteristics

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