CEVAC ND-IB-EDS K injekčná emulzia

Authorised

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

Product identification

Medicine name:
CEVAC ND-IB-EDS K injekčná emulzia

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

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

Eggdrop syndrome-1976 virus, strain B8/78, Inactivated

Target species:
Chicken

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:

Newcastle disease virus, strain La Sota, Inactivated
1.00 Protective Dose / 0.50 millilitre(s)

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

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

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Chicken**
  - All relevant tissues. 0 day

Intramuscular use:

- **Chicken**
  - 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:
Slovakia

Package description:
Available only in Slovak
Available only in Slovak
Available only in Slovak

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:
Ceva Animal Health Slovakia s.r.o.

Marketing authorisation date:
4/08/2003

Manufacturing sites for batch release:
Ceva-Phylaxia Veterinary Biologicals Co. Ltd

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

Authorisation number:
97/050/03-S

Date of authorisation status change:
23/11/2022

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

Documents

Combined File of all Documents

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