CEVAC ND-IB-IBD K vakcina A.U.V.

Authorised

- Infectious bursal disease virus, Inactivated
- Newcastle disease virus, strain La Sota, Inactivated
- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Product identification

Medicine name:
CEVAC ND-IB-IBD K vakcina A.U.V.

Active substance:
Infectious bursal disease virus, Inactivated

Newcastle disease virus, strain La Sota, Inactivated

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Target species:
Chicken (hen)

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Infectious bursal disease virus, Inactivated

3.00 log10 virus neutralising unit(s) / 1.00 unit(s)/dose

Newcastle disease virus, strain La Sota, Inactivated
4.00 log2 haemagglutination inhibiting unit(s) / 1.00 unit(s)/dose

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
6.00 log2 haemagglutination inhibiting unit(s) / 1.00 unit(s)/dose

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Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Intramuscular use:

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

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AA08

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Hungary

Package description:
Available only in Hungarian
Available only in Hungarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva-Phylaxia Zrt.

Marketing authorisation date:
25/02/2004

Manufacturing sites for batch release:
Ceva-Phylaxia Zrt.

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
25/02/2004

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