CEVAC Megamune ND-IB-EDS-SHS K vakcina A.U.V.

Authorised

- Newcastle disease virus, strain La Sota, Inactivated
- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Eggdrop syndrome-1976 virus, strain B8/78, Inactivated
- Infectious bronchitis virus, strain QX FR, Inactivated
- Turkey rhinotracheitis virus, strain TRT50, Inactivated

Product identification

Medicine name:
CEVAC Megamune ND-IB-EDS-SHS K vakcina A.U.V.

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

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

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

Infectious bronchitis virus, strain QX FR, Inactivated

Turkey rhinotracheitis virus, strain TRT50, Inactivated

Target species:
Chicken

Route of administration:
Intramuscular use

Product details

Active substance and strength:

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
4.90 log2 haemagglutination inhibiting unit(s) / 1.00 unit(s)/dose

Eggdrop syndrome-1976 virus, strain B8/78, Inactivated
7.00 log2 haemagglutination inhibiting unit(s) / 1.00 unit(s)/dose

Infectious bronchitis virus, strain QX FR, Inactivated
1.00 relative potency / 1.00 unit(s)/dose

Turkey rhinotracheitis virus, strain TRT50, Inactivated
20.00 enzyme-linked immunosorbent assay unit / 1.00 unit(s)/dose

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Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA18

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
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:
13/12/2019

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:
13/12/2019

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