

AVIGAL NRBG injekčná emulzia pre kurčatá

Authorised

- Newcastle disease virus, strain La Sota, Inactivated
- Avian reovirus, strain S1133, Inactivated
- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Infectious bursal disease virus, strain Winterfield 2512 (intermediate plus), Inactivated

Product identification

Medicine name:

AVIGAL NRBG injekčná emulzia pre kurčatá

Active substance:

Newcastle disease virus, strain La Sota, Inactivated

Avian reovirus, strain S1133, Inactivated

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Infectious bursal disease virus, strain Winterfield 2512 (intermediate plus), Inactivated

Target species:

Chicken

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Newcastle disease virus, strain La Sota, Inactivated

7.00 log₂ haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Avian reovirus, strain S1133, Inactivated

913.00 enzyme-linked immunosorbent assay unit / 0.50 millilitre(s)

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

710.00 enzyme-linked immunosorbent assay unit / 0.50 millilitre(s)

Infectious bursal disease virus, strain Winterfield 2512 (intermediate plus),
Inactivated

357.00 enzyme-linked immunosorbent assay unit / 0.50 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Chicken

- All relevant tissues. 0 day

0 dní. Pri odporazení zvierat v dobe 21 dní po vakcinácii sa odporúča odstrániť tkanivo v mieste vpichu a v jeho okolí

Subcutaneous use:

-

Chicken

- All relevant tissues. 0 day

0 dní. Pri odporazení zvierat v dobe 21 dní po vakcinácii sa odporúča odstrániť tkanivo v mieste vpichu a v jeho okolí

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA16

Legal status of supply:

This information is not available for this product.

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 - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Pharmagal Bio spol. s r.o.

Marketing authorisation date:

13/03/2003

Manufacturing sites for batch release:

Pharmagal Bio spol. s r.o.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

97/017/03-S

Date of authorisation status change:

29/11/2022

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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