

Nobilis Multriva

Authorised

RT+IBm+ND+Gm+REOm+EDS (--) Emulsion for injection

- Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated
- Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Newcastle disease virus, strain Ulster, Inactivated
- Eggdrop syndrome-1976 virus, strain BC14, Inactivated
- Infectious bursal disease virus, strain GB02, Inactivated
- Infectious bursal disease virus, strain 89/03, Inactivated
- Avian reovirus, strain ARV-1, Inactivated
- Avian reovirus, strain ARV-4, Inactivated

Product identification

Medicine name:

Nobilis Multriva RT+IBm+ND+Gm+REOm+EDS (--) Emulsion for injection

Active substance:

Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated

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

Newcastle disease virus, strain Ulster, Inactivated

Eggdrop syndrome-1976 virus, strain BC14, Inactivated

Infectious bursal disease virus, strain GB02, Inactivated

Infectious bursal disease virus, strain 89/03, Inactivated

Avian reovirus, strain ARV-1, Inactivated

Avian reovirus, strain ARV-4, Inactivated

Target species:

Chicken

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated

Presentation_strength: ≥ 19.0 U Reference: EMEA/V/VAMF/0001 Index: 3

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

Presentation_strength: ≥ 4.8 log₂ HI Reference: EMEA/V/VAMF/0002 Index: 4

Newcastle disease virus, strain Ulster, Inactivated

Presentation_strength: ≥ 5.9 U Reference: EMEA/V/VAMF/0004 Index: 5

Eggdrop syndrome-1976 virus, strain BC14, Inactivated

Presentation_strength: ≥ 368.3 U Reference: EMEA/V/VAMF/0009 Index: 6

Infectious bursal disease virus, strain GB02, Inactivated

Presentation_strength: ≥ 100.9 U1 Reference: EMEA/V/VAMF/0005 Index: 7

Infectious bursal disease virus, strain 89/03, Inactivated

Presentation_strength: ≥ 88.6 U Reference: EMEA/V/VAMF/0006 Index: 8

Avian reovirus, strain ARV-1, Inactivated

Presentation_strength: ≥ 11.5 U Reference: EMEA/V/VAMF/0007 Index: 9

Avian reovirus, strain ARV-4, Inactivated

Presentation_strength: ≥ 11.4 U1 Reference: EMEA/V/VAMF/0008 Index: 10

Pharmaceutical form:

Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA24

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Bottle (PET), Package_size:1 bottle, Content:600 ml (2000 doses)

Packaging:Bottle (PET), Package_size:1 bottle, Content:300 ml (1000 doses)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - known active substance (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

6/05/2024

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

6/05/2024

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

Documents

Combined File of all Documents

English (PDF)

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