

NOBILIS IB+G+ND emulzija za injiciranje za plemenske piščance

Not
authorised

- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Newcastle disease virus, strain Clone 30, Inactivated
- Infectious bursal disease virus, strain D78, Inactivated

Product identification

Medicine name:

NOBILIS IB+G+ND emulzija za injiciranje za plemenske piščance

Active substance:

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Newcastle disease virus, strain Clone 30, Inactivated

Infectious bursal disease virus, strain D78, Inactivated

Target species:

Chicken (for reproduction)

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

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

Newcastle disease virus, strain Clone 30, Inactivated
50.00 50% Protective Dose / 0.50 millilitre(s)

Infectious bursal disease virus, strain D78, Inactivated
14.50 log₂ virus neutralising unit(s) / 0.50 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Chicken (for reproduction)**
- Not specified. 0 day

Subcutaneous use:

- **Chicken (for reproduction)**
- Not specified. 0 day
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA08

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Slovenia

Package description:

Available only in [Slovenian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

26/06/2000

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:

NP/V/0217/001

Date of authorisation status change:

12/10/2023

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.