Nobilis IBmulti+ND+EDS Emulsion for injection (water-in-oil)

Not authorised

- Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Avian infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
- Egg drop syndrome '76 virus, strain BC14, Inactivated
- Newcastle disease virus, strain Clone 30, Inactivated

Product identification

Medicine name:
Nobilis IBmulti+ND+EDS Emulsion for injection (water-in-oil)
Nobilis IB multi + ND + EDS Emulsie voor injectie
Nobilis IB multi + ND + EDS Emulsion injectable
Nobilis IB multi + ND + EDS Emulsion zur Injektion

Active substance:
Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
Avian infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
Egg drop syndrome '76 virus, strain BC14, Inactivated
Newcastle disease virus, strain Clone 30, Inactivated

Target species:
Chicken (for reproduction)

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
2.46 virus neutralising unit(s) / 0.50 millilitre(s)

Avian infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
2.00 virus neutralising unit(s) / 0.50 millilitre(s)

Egg drop syndrome '76 virus, strain BC14, Inactivated
2.70 log10 haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Newcastle disease virus, strain Clone 30, Inactivated
2.00 log10 haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Chicken (for reproduction)**

  - Egg. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA13

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Belgium

Package description:

(ID1): 1 Box with 1 Bottle (PolyEthylene TerePhthalate) with 250 millilitre(s) (500 ID)
(ID2): 1 Box with 1 Bottle (PolyEthylene TerePhthalate) with 500 millilitre(s) (1000 ID)

(ID4): 1 Box with 1 Bottle (Glass) with 500 millilitre(s) (1000 ID)
(ID3): 1 Box with 1 Bottle (Glass) with 250 millilitre(s) (500 ID)

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:
Intervet International B.V.

Marketing authorisation date:
19/12/2005

Manufacturing sites for batch release:
INTERVET INTERNATIONAL B.V.

Responsible authority:
Federal Agency For Medicines And Health Products

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

Date of authorisation status change:
25/07/2023

Reference member state:
Germany

Procedure number:
DE/V/0225/001

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 9/08/2022
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Package Leaflet

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