Nobilis Reo Inac Emulsie voor injectie

Authorised

- Avian reovirus, strain 1733, Inactivated
- Avian reovirus, strain 2408, Inactivated

Product identification

Medicine name:
Nobilis Reo Inac Emulsie voor injectie

Active substance:
Avian reovirus, strain 1733, Inactivated
Avian reovirus, strain 2408, Inactivated

Target species:
Chicken (for reproduction)

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Avian reovirus, strain 1733, Inactivated
7.40 enzyme-linked immunosorbent assay unit / 0.50 millilitre(s)

Avian reovirus, strain 2408, Inactivated
7.40 enzyme-linked immunosorbent assay unit / 0.50 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Chicken (for reproduction)**
  - Meat and offal. 0 day
  - Egg. 0 day

Subcutaneous use:

- **Chicken (for reproduction)**
  - Meat and offal. 0 day
  - Egg. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD07

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
Nobilis Reo Inac 1 glass Vial with 1000 doses of Emulsion for injection
Nobilis Reo Inac 1 glass Vial with 500 doses of Emulsion for injection
Nobilis Reo Inac 1 PET Vial with 1000 doses of Emulsion for injection

Nobilis Reo Inac 1 PET Vial with 500 doses of Emulsion for injection

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
22/03/1999

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:
15/04/2024

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

Documents

Package Leaflet

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Labelling

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

Summary of Product Characteristics

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