BIGOPEST EMULSION INJECTABLE

Authorised

- Infectious bursal disease virus, strain VNJO, Inactivated
- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Newcastle disease virus, strain Ulster, Inactivated

Product identification

Medicine name:
BIGOPEST EMULSION INJECTABLE

Active substance:
Infectious bursal disease virus, strain VNJO, Inactivated

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Newcastle disease virus, strain Ulster, Inactivated

Target species:
Chicken

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Infectious bursal disease virus, strain VNJO, Inactivated

5.00 50% Protective Dose / 1.00 Dose

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
80.00 haemagglutination inhibiting unit(s) / 1.00 Dose

Newcastle disease virus, strain Ulster, Inactivated
16.00 50% Protective Dose / 1.00 Dose

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Chicken**
  - Eggs. 0 day

Subcutaneous use:

- **Chicken**
  - Eggs. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AA06

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

Authorisation status:
Valid

Authorised in:
France

Package description:
Available only in French
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
26/02/1990

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/1582974 6/1990

Date of authorisation status change:
18/12/2023

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

Documents

Summary of Product Characteristics

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Package Leaflet and Labelling

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