IZOVAC ND IB IBD REO

Authorised

- Infectious bursal disease virus, strain Winterfield 2512 (intermediate plus), Inactivated
- Avian reovirus, strain S1133, Inactivated
- Newcastle disease virus, strain Ulster, Inactivated
- Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Product identification

Medicine name:
IZOVAC ND IB IBD REO

Active substance:
Infectious bursal disease virus, strain Winterfield 2512 (intermediate plus), Inactivated

Avian reovirus, strain S1133, Inactivated

Newcastle disease virus, strain Ulster, Inactivated

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

Target species:
Chicken (layer hen)
Chicken (for reproduction)

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

Infectious bursal disease virus, strain Winterfield 2512 (intermediate plus), Inactivated
1.00 relative potency / 1.00 Dose

Avian reovirus, strain S1133, Inactivated
1.00 relative potency / 1.00 Dose

Newcastle disease virus, strain Ulster, Inactivated
16.00 haemagglutination inhibiting unit(s) / 1.00 Dose

Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
64.00 haemagglutination inhibiting unit(s) / 1.00 Dose

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Chicken (layer hen)**
  - Meat and offal. 0 day
  - Eggs. 0 day
- **Chicken (for reproduction)**
  - Meat and offal. 0 day

Subcutaneous use:

- **Chicken (layer hen)**
  - Meat and offal. 0 day
  - Eggs. 0 day
- **Chicken (for reproduction)**

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AA16

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

Authorisation status:
Valid

Authorised in:
Italy

Package description:
Available only in Italian
Available only in Italian

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:
Izo S.r.l.

Marketing authorisation date:
5/06/2015

Manufacturing sites for batch release:
Izo S.r.l.

Responsible authority:
Ministry Of Health

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

Date of authorisation status change:
1/02/2016

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

Documents

Combined File of all Documents

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