IZOVAC ND IB IBD

Authorised

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

Product identification

Medicine name:
IZOVAC ND IB IBD

Active substance:
Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Newcastle disease virus, strain Ulster, Inactivated

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

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

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

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

Newcastle disease virus, strain Ulster, Inactivated
16.00 haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Infectious bursal disease virus, strain Winterfield 2512 (intermediate plus), Inactivated
1.00 relative potency / 0.50 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Chicken (layer hen)**
  - Eggs. 0 day

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

Subcutaneous use:

- **Chicken (layer hen)**
  - Eggs. 0 day

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA08

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:
25/05/2017

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:
25/05/2017

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

Documents

Combined File of all Documents

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