

IZOVAC ND-EDS-IB – Emulsione iniettabile per polli

Authorised

- Infectious bronchitis virus, strain D1466, Inactivated
- Infectious bronchitis virus, strain D274, Inactivated
- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Eggdrop syndrome-1976 virus, strain V127, Inactivated
- Newcastle disease virus, strain Ulster, Inactivated

Product identification

Medicine name:

IZOVAC ND-EDS-IB – Emulsione iniettabile per polli

Active substance:

Infectious bronchitis virus, strain D1466, Inactivated

Infectious bronchitis virus, strain D274, Inactivated

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Eggdrop syndrome-1976 virus, strain V127, Inactivated

Newcastle disease virus, strain Ulster, Inactivated

Target species:

Chicken (layer hen)

Chicken (for reproduction)

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Infectious bronchitis virus, strain D1466, Inactivated

1.00 European Pharmacopoeia Unit(s) / 0.50 millilitre(s)

Infectious bronchitis virus, strain D274, Inactivated

1.00 European Pharmacopoeia Unit(s) / 0.50 millilitre(s)

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

1.00 European Pharmacopoeia Unit(s) / 0.50 millilitre(s)

Eggdrop syndrome-1976 virus, strain V127, Inactivated

1.00 European Pharmacopoeia Unit(s) / 0.50 millilitre(s)

Newcastle disease virus, strain Ulster, Inactivated

50.00 50% Protective Dose / 0.50 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

-

Chicken (layer hen)

- Meat and offal. 0 day

- Egg. 0 day

-

Chicken (for reproduction)

- Meat and offal. 0 day

- Egg. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA13

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

Available only in Italian

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:

Izo S.r.l.

Marketing authorisation date:

1/02/1993

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:

31/12/2007

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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