OLVAC A+B Emulsione iniettabile per polli

Authorised

- Newcastle disease virus, Inactivated
- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Eggdrop syndrome-1976 virus, strain V127, Inactivated
- Infectious bronchitis virus, strain NEV14, Inactivated
- Infectious bronchitis virus, strain NEV24, Inactivated

Product identification

Medicine name:
OLVAC A+B Emulsione iniettabile per polli

Active substance:
Newcastle disease virus, Inactivated
Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
Eggdrop syndrome-1976 virus, strain V127, Inactivated
Infectious bronchitis virus, strain NEV14, Inactivated
Infectious bronchitis virus, strain NEV24, Inactivated

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

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Newcastle disease virus, Inactivated
8.50 log 10 50% embryo infective dose / 0.50 Dose

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
7.50 log 10 50% embryo infective dose / 0.50 Dose

Eggdrop syndrome-1976 virus, strain V127, Inactivated
7.50 log 10 50% embryo infective dose / 0.50 Dose

Infectious bronchitis virus, strain NEV14, Inactivated
7.50 log 10 50% embryo infective dose / 0.50 Dose

Infectious bronchitis virus, strain NEV24, Inactivated
7.50 log 10 50% embryo infective dose / 0.50 Dose

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

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

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA13

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:
Italy

Package description:
Available only in Italian
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:
Fatro S.p.A.

Marketing authorisation date:
28/02/1990

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Ministry Of Health

Authorisation number:
101816

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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