

OLVAC A +B+MG Vaccino inattivato contro la Pseudopeste aviare, la Bronchite infettiva, la sindrome del calo di deposizione (EDS'76) e l'infezione da Mycoplasma gallisepticum

Authorised

- Newcastle disease virus, Inactivated
- Avian infectious bronchitis virus, Inactivated
- Eggdrop syndrome-1976 virus, Inactivated
- Mycoplasma gallisepticum, Inactivated

Product identification

Medicine name:

OLVAC A +B+MG Vaccino inattivato contro la Pseudopeste aviare, la Bronchite infettiva, la sindrome del calo di deposizione (EDS'76) e l'infezione da Mycoplasma gallisepticum

Active substance:

Newcastle disease virus, Inactivated

Avian infectious bronchitis virus, Inactivated

Eggdrop syndrome-1976 virus, Inactivated

Mycoplasma gallisepticum, Inactivated

Target species:

Chicken (layer hen)

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Newcastle disease virus, Inactivated

50.00 50% Protective Dose / 1.00 Dose

Avian infectious bronchitis virus, Inactivated

7.50 log 10 50% embryo infective dose / 1.00 Dose

Eggdrop syndrome-1976 virus, Inactivated

1000.00 haemagglutinating units / 1.00 Dose

Mycoplasma gallisepticum, Inactivated

30.00 log10 colony forming unit(s) / 1.00 Dose

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:**Subcutaneous use:**

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Chicken (layer hen)

- Egg. 0 day

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AL

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](#)

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:

25/10/2001

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

Ministry Of Health

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

25/10/2001

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

Documents

Summary of Product Characteristics

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