OLVAC A+B+HG

Authorised

- Avibacterium paragallinarum, serotype C, strain Modesto, Inactivated
- Newcastle disease virus, Inactivated
- Avian infectious bronchitis virus, Inactivated
- Eggdrop syndrome-1976 virus, Inactivated
- Avibacterium paragallinarum, serotype A, strain W, Inactivated

Product identification

Medicine name:
OLVAC A+B+HG

Active substance:
Avibacterium paragallinarum, serotype C, strain Modesto, Inactivated

Newcastle disease virus, Inactivated

Avian infectious bronchitis virus, Inactivated

Eggdrop syndrome-1976 virus, Inactivated

Avibacterium paragallinarum, serotype A, strain W, Inactivated

Target species:
Chicken (layer hen)

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Avibacterium paragallinarum, serotype C, strain Modesto, Inactivated
3.00 billion colony forming units / 0.50 millilitre(s)

Newcastle disease virus, Inactivated
50.00 50% Protective Dose / 0.50 millilitre(s)

Avian infectious bronchitis virus, Inactivated
7.50 log 10 50% embryo infective dose / 0.50 millilitre(s)

Eggdrop syndrome-1976 virus, Inactivated
1000.00 haemagglutinating units / 0.50 millilitre(s)

Avibacterium paragallinarum, serotype A, strain W, Inactivated
3.00 billion colony forming units / 0.50 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AL

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
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

Combined File of all Documents

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