NOBILIS IB MULTI+ND+EDS ΕΝΕΣΙΜΟ ΓΑΛΑΚΤΩΜΑ

Authorised

- Eggdrop syndrome-1976 virus, Inactivated
- Newcastle disease virus, Inactivated
- Avian infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
- Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Product identification

Medicine name:
NOBILIS IB MULTI+ND+EDS ΕΝΕΣΙΜΟ ΓΑΛΑΚΤΩΜΑ

Active substance:
Eggdrop syndrome-1976 virus, Inactivated

Newcastle disease virus, Inactivated

Avian infectious bronchitis virus, type D274/D207, strain 249g, Inactivated

Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Target species:
Chicken

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

Eggdrop syndrome-1976 virus, Inactivated
6.50 log2 virus neutralising unit(s) / 0.50 millilitre(s)

Newcastle disease virus, Inactivated
4.00 log2 haemagglutination inhibiting unit(s) / 0.01 millilitre(s)

Avian infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
4.00 log2 virus neutralising unit(s) / 0.50 millilitre(s)

Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
4.00 log2 virus neutralising unit(s) / 0.50 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Chicken**
  - Not applicable. no withdrawal period

Subcutaneous use:

- **Chicken**
  - Not applicable. no withdrawal period

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:

Greece

Package description:
Available only in Greek
Available only in Greek

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone)

Marketing authorisation holder:
Intervet Hellas A.E.

Marketing authorisation date:
11/10/1999

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
National Organization For Medicines

Authorisation number:
34517/12-10-1999/K-0126401

Date of authorisation status change:
14/10/2007

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