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BIO NEW EDS + IB, emulsione iniettabile per polli

Authorised

- Newcastle disease virus, strain Ulster 2C, Inactivated
- Eggdrop syndrome-1976 virus, strain McFerran 127, Inactivated
- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Product identification

Medicine name:
BIO NEW EDS + IB, emulsione iniettabile per polli

Active substance:
Newcastle disease virus, strain Ulster 2C, Inactivated

Eggdrop syndrome-1976 virus, strain McFerran 127, Inactivated

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Target species:
Chicken (pullet)

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Newcastle disease virus, strain Ulster 2C, Inactivated
4.00 log2 haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Eggdrop syndrome-1976 virus, strain McFerran 127, Inactivated
7.32 log2 haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
6.00 log2 haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Chicken (pullet)**
  - Egg. 0 day
  - Meat and offal. 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

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:
Boehringer Ingelheim Animal Health Italia S.p.A. In Breve Boehringer Ingelheim Ah It S.p.A.

Marketing authorisation date:
19/11/1993

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health Italia S.p.A. In Breve Boehringer Ingelheim Ah It S.p.A.

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
19/11/2008

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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