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NOBILIS E.COLI INAC

Authorised

- Escherichia coli, fimbrial adhesin F11
- Escherichia coli, flagellar toxin

Product identification

Medicine name:
NOBILIS E.COLI INAC

Active substance:
Escherichia coli, fimbrial adhesin F11

Escherichia coli, flagellar toxin

Target species:
Chicken

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Escherichia coli, fimbrial adhesin F11
100.00 microgram(s) / 0.50 millilitre(s)

Escherichia coli, flagellar toxin
100.00 microgram(s) / 0.50 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Chicken**
  - Meat and offal. 35 day
  - Egg. 0 day

Subcutaneous use:

- **Chicken**
  - Meat and offal. 35 day
  - Egg. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AB05

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
250 ml (500 doses) in one glass type II vial closed with a nitryl rubber stopper and sealed with a coded aluminium cap in a cardboard box.
500 ml (1000 doses) in one glass type II vial closed with a nitryl rubber stopper and sealed with a coded aluminium cap in a cardboard box.

500 ml (1000 doses) in one PET vial closed with a nitryl rubber stopper and sealed with a coded aluminium cap in a cardboard box.
250 ml (500 doses) in one PET vial closed with a nitryl rubber stopper and sealed with a coded aluminium cap in a cardboard box.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Intervet Deutschland GmbH

Marketing authorisation date:
25/01/1995

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

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
283a/93

Date of authorisation status change:
12/10/2010

Reference member state:
Netherlands

Procedure number:
NL/V/0017/001

Concerned member states:
Austria Belgium France Germany Greece Ireland Italy Portugal Spain

United Kingdom (Northern Ireland)

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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