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Nobilis CAV P4

Authorised

- Chicken anaemia virus, strain 26P4, Live

Product identification

Medicine name:
Nobilis CAV P4

Active substance:
Chicken anaemia virus, strain 26P4, Live

Target species:
Chicken

Route of administration:
Wing-web-stab use
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Chicken anaemia virus, strain 26P4, Live
3.00 log10 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate for suspension for injection

Withdrawal period by route of administration:

Wing-web-stab use:

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

Intramuscular use:

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

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD04

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

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Authorisation status:
Valid

Authorised in:
Denmark

Available in:
Denmark

Package description:
Cardboard box with 1 x 1000 doses Lyophilisate: 10 ml glass vial (type I) with halogenobutyl rubber stopper and coded aluminium capsule

Cardboard box with 1 x 13 ml solvent Unisolve: 20 ml glass vial (type II) with halogenobutyl rubber stopper and coded aluminium capsule
Cardboard box with 1 x 200 ml solvent Dilavia: 250 ml glass (type II) or PET vial with halogenobutyl rubber stopper and coded aluminium capsule
Cardboard box with 10 x 1000 doses of vaccine - Lyophilisate: 10 ml glass vial (type I) with halogenobutyl rubber stopper and coded aluminium capsule
Cardboard box with 10 x 13 ml solvent Unisolve: 20 ml glass vial (type II) with halogenobutyl rubber stopper and coded aluminium capsule.
Cardboard box with 10 x 200 ml solvent Dilavia: 250 ml glass (type II) or PET vial with halogenobutyl rubber stopper and coded aluminium capsule

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
27/01/1997

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

Responsible authority:
Danish Medicines Agency

Authorisation number:
18039

Date of authorisation status change:
27/01/1997

Reference member state:
Netherlands

Procedure number:
NL/V/8659/001

Concerned member states:
Belgium Denmark Finland Germany Portugal

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 26/08/2024
Updated on: 14/03/2026
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