AVINEW NEO EFFERVESCENT TABLET FOR CHICKENS AND TURKEYS

Not authorised

- Newcastle disease virus, strain VG/GA, Live

Product identification

Medicine name:
AVINEW NEO EFFERVESCENT TABLET FOR CHICKENS AND TURKEYS

Active substance:
Newcastle disease virus, strain VG/GA, Live

Target species:
Chicken (layer hen)
Chicken (for reproduction)
Chicken (broiler)
Turkey

Route of administration:
Ocular use
Oral use
Oculonasal use

Product details

Active substance and strength:
Newcastle disease virus, strain VG/GA, Live
5.50 log10 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:
Effervescent tablet

Withdrawal period by route of administration:
Ocular use:

- **Chicken (layer hen)**
  - All relevant tissues. 0 day
- **Chicken (for reproduction)**
  - All relevant tissues. 0 day
- **Chicken (broiler)**
  - All relevant tissues. 0 day

Oral use:

- **Chicken (layer hen)**
  - All relevant tissues. 0 day
- **Chicken (for reproduction)**
  - All relevant tissues. 0 day
- **Chicken (broiler)**
  - All relevant tissues. 0 day

Oculonasal use:

- **Turkey**
  - All relevant tissues. 0 day
- **Chicken (for reproduction)**
  - All relevant tissues. 0 day
- **Chicken (broiler)**
  - All relevant tissues. 0 day
- **Chicken (layer hen)**
  - All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD06

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

Authorisation status:
Surrendered

Authorised in:
Netherlands

Package description:
Box of 10 blisters of 10 tablets of 2,000 doses
Box of 1 blister of 10 tablets of 1,000 doses
Box of 1 blister of 10 tablets of 2,000 doses
Box of 10 blisters of 10 tablets of 1,000 doses

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Animal Health Netherlands B.V.

Marketing authorisation date:
4/12/2015

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 117276

Date of authorisation status change:
22/09/2025

Reference member state:
France

Procedure number:
FR/V/0296/001

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

Documents

Combined File of all Documents

English (PDF)

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