File downloaded on 2026-11-28
Source URL: https://medicines.health.europa.eu/veterinary/en/600000042168

AVINEW

Authorised

- Newcastle disease virus, strain VG/GA-AVINEW, Live

Product identification

Medicine name:
AVINEW

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

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

Route of administration:
Ocular use
Oral use
Oculonasal use

Product details

Active substance and strength:
Newcastle disease virus, strain VG/GA-AVINEW, Live
5.50 log 10 50% embryo infective dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate for oculonasal suspension/use in drinking water

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:

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

Chicken (broiler)

- 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:
Valid

Authorised in:
France

Package description:
Box of one 1,000-dose bottle
Box of ten 2,000-dose bottles
Box of ten 1,000-dose bottles
Box of one 2,000-dose bottle

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
13/12/1999

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/8349895 2/1999

Date of authorisation status change:
13/12/2009

Reference member state:
France

Procedure number:
FR/V/0123/001

Concerned member states:
Belgium Germany Greece Luxembourg Netherlands Portugal United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.