AVINEW

Not authorised

- Newcastle disease virus, strain VG/GA, Live

Product identification

Medicine name:
AVINEW
AVINEW

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

Pharmaceutical form:

Lyophilisate for suspension

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:
Surrendered

Authorised in:
Greece

Package description:
Box of one 2,000-dose bottle
Box of ten 1,000-dose bottles
Box of ten 2,000-dose bottles
Box of one 1,000-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:
28/04/2002

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
National Organization For Medicines

Authorisation number:
49731/20-07-2007/K-0140801

Date of authorisation status change:
30/09/2025

Reference member state:
France

Procedure number:
FR/V/0123/001

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