HuveGuard NB

Authorised

- Eimeria necatrix, strain Mednec 3+8, Live
- Eimeria brunetti, strain Roybru 3+28, Live

Product identification

Medicine name:
HuveGuard NB
HuveGuard NB εναιώρημα για από του στόματος χρήση σε όρνιθες

Active substance:
Eimeria necatrix, strain Mednec 3+8, Live

Eimeria brunetti, strain Roybru 3+28, Live

Target species:
Chicken

Route of administration:
Oral use

Product details

Active substance and strength:
Eimeria necatrix, strain Mednec 3+8, Live
100.00 unit(s) / 1.00 Dose

Eimeria brunetti, strain Roybru 3+28, Live
50.00 unit(s) / 1.00 Dose

Pharmaceutical form:

Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Chicken**

  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AN01

Legal status of supply:

This information is not available for this product.

Authorisation status:

Valid

Authorised in:

Greece

Available in:

Greece

Package description:

Low density polyethylene (LDPE) vial of 30 ml with a grey butyl rubber stopper and aluminium cap. Cardboard box with 5 vials of 5000 doses.
Low density polyethylene (LDPE) vial of 30 ml with a grey butyl rubber stopper and aluminium cap. Cardboard box with 10 vials of 1000 doses.
Low density polyethylene (LDPE) vial of 30 ml with a grey butyl rubber stopper and aluminium cap. Cardboard box with 1 vial of 5000 doses
Low density polyethylene (LDPE) vial of 30 ml with a grey butyl rubber stopper and aluminium cap. Cardboard box with 1 vial of 1000 doses.
Low density polyethylene (LDPE) vial of 30 ml with a grey butyl rubber stopper and aluminium cap. Cardboard box with 5 vials of 1000 doses.
Low density polyethylene (LDPE) vial of 30 ml with a grey butyl rubber stopper and aluminium cap. Cardboard box with 10 vials of 5000 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:
HuVepharma

Marketing authorisation date:
This information is not available for this product.

Manufacturing sites for batch release:
Biovet J.S.C.

Responsible authority:
National Organization For Medicines

Authorisation number:
107716/29-11-2017/K-0231001

Date of authorisation status change:
2/06/2021

Reference member state:
Netherlands

Procedure number:
NL/V/0207/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Ireland Italy Latvia Lithuania Malta Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden 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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