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HuveGuard NB suspension for oral suspension for chickens

Authorised

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

Product identification

Medicine name:
HuveGuard NB suspension for oral suspension for chickens

Active substance:
Eimeria brunetti, strain Roybru 3+28, Live

Eimeria necatrix, strain Mednec 3+8, Live

Target species:
Chicken

Route of administration:
Oral use

Product details

Active substance and strength:
Eimeria brunetti, strain Roybru 3+28, Live
50.00 oocyst(s) / 0.03 millilitre(s)

Eimeria necatrix, strain Mednec 3+8, Live

100.00 oocyst(s) / 0.03 millilitre(s)

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:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Sweden

Package description:
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 (1 dose = 0.025 ml)
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 (1 dose = 0.025 ml)
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 (1 dose = 0.025 ml)
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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 (1 dose = 0.025 ml)

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:
15/02/2018

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
54277

Date of authorisation status change:
15/02/2018

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

Package Leaflet

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

Summary of Product Characteristics

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

Labelling

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

Combined File of all Documents