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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:
Germany

Available in:
Germany

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)
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 (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:
20/06/2016

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
PEI.V.11837.01.1

Date of authorisation status change:
14/10/2020

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)

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Documents

Combined File of all Documents

English (PDF)
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