

# 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

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**Active substance:**

Eimeria brunetti, strain Roybru 3+28, Live

Eimeria necatrix, strain Mednec 3+8, Live

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**Target species:**

Chicken

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**Route of administration:**

Oral use

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## 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)

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**Pharmaceutical form:**

Oral suspension

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**Withdrawal period by route of administration:**

**Oral use:**

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**Chicken**

- Meat and offal. 0 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI01AN01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Ireland

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**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)

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 (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 5000 doses (1 dose = 0.025 ml)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

HuVepharma

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**Marketing authorisation date:**

19/08/2016

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**Manufacturing sites for batch release:**

Biovet AD

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA10782/025/001

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**Date of authorisation status change:**

19/08/2016

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0207/001

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**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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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents