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# HuveGuard NB suspension for oral suspension for chickens

Ima dovoljenje za promet

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

## Informacije o zdravilu

**Ime zdravila:**
HuveGuard NB suspension for oral suspension for chickens

**Učinkovina:**
Na voljo samo v English
Na voljo samo v English

**Ciljne živalske vrste:**
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

**Pot uporabe:**
Peroralna uporaba

## Podatki o zdravilu

**Učinkovina / Jakost:**
Na voljo samo v English
50.00 oocyst(s) / 0.03 millilitre(s)

Na voljo samo v English
100.00 oocyst(s) / 0.03 millilitre(s)

**Farmacevtska oblika:**

Peroralna suspenzija

**Karenca glede na pot uporabe:**

**Peroralna uporaba:**

- **Chicken**
  - Meat and offal. 0 day

**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**

QI01AN01

**Pravni status za dobavo/izdajo zdravila:**

Zdravilo, ki se izdaja na veterinarski recept

**Status dovoljenja za promet z zdravilom:**

Valid

**Ima dovoljenje za promet v:**

Na voljo samo v Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

**Na voljo v:**

Spain

**Opis ovojnine zdravila:**

Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

## Dodatne informacije

**Vrsta dovoljenja:**
Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

**Pravna podlaga za izdajo dovoljenja za promet z zdravilom:**
Na voljo samo v English Italian Latvian Lithuanian Norwegian

**Imetnik dovoljenja za promet z zdravilom:**
HuVepharma

**Datum dovoljenja za promet z zdravilom:**
12/07/2016

**Proizvodna mesta, odgovorna za sproščanje serij:**
Biovet AD

**Pristojni organ:**
Spanish Agency Of Medicines And Medical Devices

**Številka dovoljenja za promet z zdravilom:**
3428 ESP

**Datum spremembe statusa dovoljenja za promet:**
25/12/2021

**Referenčna država članica:**
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

**Številka postopka:**
NL/V/0207/001

**Zadevne države članice:**

Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

## Dokumenti

Povzetek glavnih značilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

Navodilo za uporabo

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

Označevanje

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.