Tabic V.H. putojošā tablete vistām un tītariem

Ovlašten

- Newcastle disease virus, strain V.H., Live

Product identification

Naziv VMP-a:
Tabic V.H. putojošā tablete vistām un tītariem

Djelatna tvar:
Dostupno samo u English

Ciljne vrste životinja:
Dostupno samo u Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Dostupno samo u Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
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Dostupno samo u Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Slovenian Finnish Swedish Icelandic Norwegian

Način primjene:
Nebulizacijom
U vodi za piće
Intraokularno

Product details

Djelatna tvar i jačina:

Dostupno samo u English
1000000.00 50% Embryo Infective Dose / 1.00 unit(s)

Farmaceutski oblik:

Šumeća tableta

Withdrawal period by route of administration:

Nebulizacijom:

- **Chicken (for reproduction)**
  - Not specified. 0 day
- **Chicken (layer hen)**
  - Not specified. 0 day
- **Turkey**
  - Not specified. 0 day
- **Chicken (broiler)**
  - Not specified. 0 day

U vodi za piće:

- **Chicken (broiler)**
  - Not specified. 0 day
- **Chicken (for reproduction)**
  - Not specified. 0 day
-

Chicken (layer hen)
- Not specified. 0 day

•
Turkey
- Not specified. 0 day

Intraokularno:

•
Chicken (broiler)
- Not specified. 0 day

•
Chicken (for reproduction)
- Not specified. 0 day

•
Chicken (layer hen)
- Not specified. 0 day

•
Turkey
- Not specified. 0 day

Kod anatomsko-terapijsko-kemijske klasifikacije (ATKvet kôd):
QI01AD06

Pravni status opskrbe:
Dostupno samo u Czech Estonian English French Italian Latvian Lithuanian Portuguese Romanian Slovenian Finnish Swedish Icelandic Norwegian

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Status odobrenja:
Važeće

Authorised in:
Dostupno samo u Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Available in:

Latvia

Opis paketa:
Dostupno samo u Latvian
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Dostupno samo u Latvian

Additional information

Entitlement type:
Marketing Authorisation

Pravna osnova za odobrenje proizvoda:
Dostupno samo u English Italian Latvian Norwegian

Nositelj odobrenja za stavljanje u promet:
Phibro Animal Health (Poland) Sp. z o.o.

Marketing authorisation date:
16/12/2010

Mjesta proizvodnje za izdavanje serije:
Synoptis Industrial Sp. z o.o.

Odgovorno tijelo:
Food And Veterinary Service

Broj autorizacije:
V/NRP/10/0032

Datum promjene statusa odobrenja:
16/12/2010

Za izvješća o sumnjama na nuspojave veterinarskog lijeka molimo vas otiđite na www.adrreports.eu/vet

Documents

Combined File of all Documents

Ovaj dokument ne postoji na ovom jeziku (hrvatski). U nastavku ga možete pronaći na drugom jeziku.

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