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Nobilis RT+IBmulti+ND+EDS emulsion for injection for chickens

Ima dovoljenje za promet

- Newcastle disease virus, strain Clone 30, Inactivated
- Eggdrop syndrome-1976 virus, strain BC14, Inactivated
- Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated
- Infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Informacije o zdravilu

Ime zdravila:
Nobilis RT+IBmulti+ND+EDS emulsion for injection for chickens

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Učinkovina:
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

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:
intramuskularna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
50.00 50% Protective Dose / 0.50 millilitre(s)

Na voljo samo v English
6.50 log2 haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Na voljo samo v English
9.50 log2 enzyme-linked immunosorbent assay unit(s) / 0.50 millilitre(s)

Na voljo samo v English
4.00 log2 virus neutralising unit(s) / 0.50 millilitre(s)

Na voljo samo v English
5.50 log2 virus neutralising unit(s) / 0.50 millilitre(s)

Farmacevtska oblika:
Emulzija za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

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

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI01AA18

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 Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Na voljo v:
Ireland

Opis ovojnine zdravila:
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 Norwegian

Imetnik dovoljenja za promet z zdravilom:
Intervet (Ireland) Limited

Datum dovoljenja za promet z zdravilom:
3/11/2004

Proizvodna mesta, odgovorna za sproščanje serij:
Intervet International B.V.

Pristojni organ:
Health Products Regulatory Authority

Številka dovoljenja za promet z zdravilom:
VPA10996/181/001

Datum spremembe statusa dovoljenja za promet:
3/11/2004

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

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Številka postopka:
DE/V/0209/001

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

Combined File of all Documents

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