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Ultifend ND IBD (--) - Concentrate and solvent for suspension for injection

Údaraithe

- Turkey herpesvirus, strain rHVT/ND/IBD (cell-associated), encoding fusion protein gene of Newcastle disease virus and VP2 protein gene of Infectious bursal disease virus, Live

Sainaithint táirge

Ainm an chógais:
Ultifend ND IBD (--) - Concentrate and solvent for suspension for injection

Substaint ghníomhach:
Ar fáil ach amháin i English

Speiceas:
Ar fáil ach amháin i Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Ar fáil ach amháin i Bulgarian Spanish Czech Danish Estonian English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

Bealach riartha:
San ubh shuthaithe
Úsáid fho-chraicneach

Sonraí an táirge

Substaint agus neart gníomhach:
Ar fáil ach amháin i English
Presentation_strength:4000 PFU/dose - 12000 PFU/dose Index:0

Foirm chógaisíochta:
Tiúchán agus tuaslagóir le haghaidh fuaidreáin le haghaidh insteallta

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Tréimhse tarraingthe siar de réir an bhealaigh riaracháin:
San ubh shuthaithe:

- **Chicken**
  - Not applicable. 0 day Zero days

Úsáid fho-chraicneach:

- **Chicken**
  - Not applicable. 0 day Zero days

An cód tréidliachta ceimiceach teiripeach anatamaíoch (Cód ATCvet):
QI01AD16

Stádas dlí an tsoláthair:
Ar fáil ach amháin i Czech Estonian English French Italian Latvian Lithuanian Portuguese Romanian Slovenian Finnish Swedish Icelandic Norwegian

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Stádas údaraithe:
Valid

Údaraithe i:
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Tuairisc ar an bpacáiste:

Ar fáil ach amháin i English
Ar fáil ach amháin i English
Ar fáil ach amháin i English

Faisnéis bhreise

Cineál teidlíochta:
Ar fáil ach amháin i English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Bunús dlí maidir le húdarú an táirge:
Ar fáil ach amháin i English Italian Latvian Lithuanian Norwegian

Sealbhóir an údaraithe margaíochta:
Ceva-Phylaxia Zrt.

Dáta Údaraithe Margaíochta:
20/04/2021

Láithreacha monaraíochta um eisiúint bhaisce:
Ceva-Phylaxia Zrt.

An t-údarás atá freagrach:
European Commission

Uimhir an údaraithe:
Níl an fhaisnéis seo ar fáil don táirge seo.

Athrú stádais maidir leis an dáta údaraithe:
20/04/2021

Tuairiscí maidir le fo-iarmhairtí amhrasta: www.adrreports.eu/vet

Doiciméid

Combined File of all Documents

Níl an doiciméad seo ann sa teanga seo (Gaeilge). Is féidir leat é a fháil i dteanga eile thíos.