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Chanaxin 100 mg/ml - Solution for injection

Údaraithe

- Tulathromycin

Sainaithint táirge

Ainm an chógais:
Chanaxin 100 mg/ml - Solution 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 Slovenian Finnish Swedish Icelandic Norwegian
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 German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

Bealach riartha:
Úsáid ionmhatánach
Úsáid fho-chraicneach

Sonraí an táirge

Substaint agus neart gníomhach:

Ar fáil ach amháin i English
100.00 milligram(s) / 1.00 Fial

Foirm chógaisíochta:

Tuaslagán le haghaidh insteallta

Tréimhse tarraingthe siar de réir an bhealaigh riaracháin:

Úsáid ionmhatánach:

- **Cattle**
  - Meat and offal. 22 day — 22 days
- **Sheep**
  - Meat and offal. 16 day — 16 days
- **Pig**
  - Meat and offal. 13 day — 13 days

Úsáid fho-chraicneach:

- **Cattle**
  - Meat and offal. 22 day — 22 days
- **Sheep**
  - Meat and offal. 16 day — 16 days
- **Pig**

- Meat and offal. 13 day 13 days

An cód tréidliachta ceimiceach teiripeach anatamaíoch (Cód ATCvet):
QJ01FA94

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

Stádas údaraithe:
Valid

Údaraithe i:
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Ar fáil ach amháin i Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

Tuairisc ar an bpacáiste:

Ar fáil ach amháin i English

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 Norwegian

Sealbhóir an údaraithe margaíochta:

Chanelle Pharmaceuticals Manufacturing Limited

Dáta Údaraithe Margaíochta:

19/04/2022

Láithreacha monaraíochta um eisiúint bhaisce:
Chanelle Pharmaceuticals Manufacturing Limited

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:
19/04/2022

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.

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