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Lexylan 180 mg/ml - Suspension for injection

Údaraithe

- Cefalexin

Sainaithint táirge

Ainm an chógais:
Lexylan 180 mg/ml - 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 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
180.00 milligram(s) / 1.00 millilitre(s)

Foirm chógaisíochta:
Fuaidreán le haghaidh insteallta

Tréimhse tarraingthe siar de réir an bhealaigh riaracháin:
Úsáid ionmhatánach:

- **Cattle**
  - Meat and offal. 12 day — 12 Days
  - Milk. 0 hour — Zero Hours

Úsáid fho-chraicneach:

- **Cattle**
  - Meat and offal. 12 day — 12 Days
  - Milk. 0 hour — Zero Hours

An cód tréidliachta ceimiceach teiripeach anatamaíoch (Cód ATCvet):
QJ01DB01

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:

Ar fáil ach amháin i Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Ar fáil ach amháin i Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

Ar fáil i:
Austria , Belgium , Denmark , France , Germany , Ireland , Italy , Luxembourg , Netherlands , Spain

Tuairisc ar an bpacáiste:
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 French Italian Latvian Lithuanian Norwegian

Sealbhóir an údaraithe margaíochta:
Emdoka

Dáta Údaraithe Margaíochta:
8/04/2024

Láithreacha monaraíochta um eisiúint bhaisce:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

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:
8/04/2024

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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