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Bupredine Multidose 0.3 mg/ml Solution for Injection for Dogs, Cats and Horses

Údaraithe

- Buprenorphine hydrochloride

Sainaithint táirge

Ainm an chógais:
Bupredine Multidose 0.3 mg/ml Solution for Injection for Dogs, Cats and Horses

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 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 infhéitheach

Sonraí an táirge

Substaint agus neart gníomhach:
Ar fáil ach amháin i English
0.32 milligram(s) / 1.00 millilitre(s)

Foirm chógaisíochta:
Tuaslagán le haghaidh insteallta

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

- **Horse**
  - All relevant tissues. no withdrawal period

Not authorised for use in horses intended for human consumption.

An cód tréidliachta ceimiceach teiripeach anatamaíoch (Cód ATCvet):
QN02AE01

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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Tuairisc ar an bpacáiste:
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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:
Le Vet. Beheer B.V.

Dáta Údaraithe Margaíochta:
6/03/2016

Láithreacha monaraíochta um eisiúint bhaisce:
Produlab Pharma B.V.

An t-údarás atá freagrach:
National Organization For Medicines

Uimhir an údaraithe:
35406/06-04-2022/K-0212001

Athrú stádais maidir leis an dáta údaraithe:
5/04/2022

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

Uimhir an nóis imeachta:
NL/V/0314/001

Ballstáit lena mbaineann:

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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.