Buprecare Multidose 0.3 mg/ml Solution for Injection for Dogs and Cats

Údaraithe

- Buprenorphine hydrochloride

Product identification

Ainm an chógais:
Buprecare Multidose 0.3 mg/ml Solution for Injection for Dogs and Cats
Buprecare Multidose 0,3 mg/ml Injektionslösung für Hunde und Katzen

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

Bealach riartha:
Úsáid ionmhatánach
Úsáid infhéitheach

Product details

Substaint ghníomhach / Láidreacht:

Ar fáil ach amháin i English
0.32 milligram(s) / 1.00 millilitre(s)

Foirm chógaisíochta:

Tuaslagán le haghaidh insteallta

Withdrawal period by route of administration:

Úsáid ionmhatánach:

- **Dog**
- **Cat**

Úsáid infhéitheach:

- **Dog**
- **Cat**

An cód tréidliachta ceimiceach teiripeach anatamaíoch (Cód ATCvet):

QN02AE01

Stádas dlí an tsoláthair:

Ar fáil ach amháin i Czech Estonian English French Italian Latvian Portuguese Slovenian Finnish Swedish Icelandic Norwegian

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Stádas údaraithe:

Valid

Authorised in:

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:

Ar fáil ach amháin i English

Additional information

Entitlement type:

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:
Ecuphar

Marketing authorisation date:
5/11/2011

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

An t-údarás atá freagrach:
Federal Office Of Consumer Protection And Food Safety

Uimhir údaraithe:
401450.00.00

Athrú stádais maidir leis an dáta údaraithe:
31/01/2017

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

Uimhir an nóis imeachta:
IE/V/0453/002

Ballstáit lena mbaineann:
Ar fáil ach amháin i Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Achoimre ar shaintréithe an táirge

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