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Ketexx 100 mg/ml solution for injection

Údaraithe

- Ketamine hydrochloride

Sainaithint táirge

Ainm an chógais:
Ketexx 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
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Bealach riartha:
Úsáid ionmhatánach
Úsáid infhéitheach
Úsáid inpheireatóinéim

Sonraí an táirge

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

Foirm chógaisíochta:
Tuaslagán le haghaidh insteallta

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

- **Cattle**

- Meat and offal. 1 day

- Milk. 0 day

•

Rabbit (exclusively kept as pet)

- All relevant tissues. no withdrawal period

Not authorised for use in rabbits for human consumption.

•

Horse

- Meat and offal. 1 day

- Milk. 0 day

Úsáid infhéitheach:

•

Cattle

- Meat and offal. 1 day

- Milk. 0 day

•

Rabbit (exclusively kept as pet)

- All relevant tissues. no withdrawal period

Not authorised for use in rabbits for human consumption.

•

Horse

- Meat and offal. 1 day

- Milk. 0 day

•

Sheep

- Meat and offal. 1 day

- Milk. 0 day

•

Goat

- Meat and offal. 1 day

- Milk. 0 day

Úsáid inpheireatóinéim:

- **Rabbit (exclusively kept as pet)**
  - All relevant tissues. no withdrawal period

Not authorised for use in rabbits for human consumption.

An cód tréidliachta ceimiceach teiripeach anatamaíoch (Cód ATCvet):
QN01AX03

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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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:
Alfasan Nederland B.V.

Dáta Údaraithe Margaíochta:
20/05/2022

Láithreacha monaraíochta um eisiúint bhaisce:
Alfasan Nederland B.V.

An t-údarás atá freagrach:
Health Products Regulatory Authority

Uimhir an údaraithe:
VPA10980/016/001

Athrú stádais maidir leis an dáta údaraithe:
20/05/2022

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

Uimhir an nóis imeachta:
FR/V/0435/001

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.

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