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Posatex 8.5 mg/ml orbifloxacin + 0.9 mg/ml mometasone furoate + 0.9 mg/ml posaconazole - Ear drops, suspension

Údaraithe

- Mometasone furoate
- Posaconazole
- Orbifloxacin

Sainaithint táirge

Ainm an chógais:
Posatex 8.5 mg/ml orbifloxacin + 0.9 mg/ml mometasone furoate + 0.9 mg/ml posaconazole - Ear drops, suspension

Substaint ghníomhach:
Ar fáil ach amháin i English
Ar fáil ach amháin i English
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

Bealach riartha:
Úsáid chluaisíneach

Sonraí an táirge

Substaint agus neart gníomhach:

Ar fáil ach amháin i English
0.86 milligram(s) / 1.00 Buidéal

Ar fáil ach amháin i English
0.86 milligram(s) / 1.00 Buidéal

Ar fáil ach amháin i English
8.55 milligram(s) / 1.00 Buidéal

Foirm chógaisíochta:

Braonta cluaise, fuaidreán

An cód tréidliachta ceimiceach teiripeach anatamaíoch (Cód ATCvet):

QS02CA91

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 i:

Austria , Belgium , Bulgaria , Cyprus , Czechia , France , Germany , Greece , Hungary , Ireland , Italy , Luxembourg , Norway , Poland , Portugal , Romania , Slovakia , Spain , United Kingdom (Northern Ireland)

Tuairisc ar an bpacáiste:

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

Sealbhóir an údaraithe margaíochta:
Intervet International B.V.

Dáta Údaraithe Margaíochta:
23/06/2008

Láithreacha monaraíochta um eisiúint bhaisce:
Vet Pharma Friesoythe GmbH

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:
23/06/2008

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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