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Cardalis 5 mg benazepril hydrochloride + 40 mg spironolactone - Chewable Tablet

Údaraithe

- Benazepril hydrochloride
- Spironolactone

Sainaithint táirge

Ainm an chógais:
Cardalis 5 mg benazepril hydrochloride + 40 mg spironolactone - Chewable Tablet

Substaint ghníomhach:
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 béil

Sonraí an táirge

Substaint agus neart gníomhach:

Ar fáil ach amháin i English
5.00 milligram(s) / 1.00 Táibléad

Ar fáil ach amháin i English
50.00 milligram(s) / 1.00 Táibléad

Foirm chógaisíochta:

Capsúl ionchoganta

An cód tréidliachta ceimiceach teiripeach anatamaíoch (Cód ATCvet):

QC09BA07

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 , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

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

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Bunús dlí maidir le húdarú an táirge:
Ar fáil ach amháin i English Italian Latvian Norwegian

Sealbhóir an údaraithe margaíochta:
Ceva Sante Animale

Dáta Údaraithe Margaíochta:
23/07/2012

Láithreacha monaraíochta um eisiúint bhaisce:
Catalent Germany Schorndorf GmbH
Ceva Sante Animale
Ceva Sante Animale

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:
23/07/2012

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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