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ABCEDYL GA SOLUTION BUVABLE

Ima dovoljenje za promet

- PYROGENIUM C7
- Virola sebifera C3
- Atropa bella-donna C5
- ECHINACEA C3
- Calcium sulfuricum C7
- ACIDUM SILICICUM C7
- HEPAR SULFURIS C7

Informacije o zdravilu

Ime zdravila:
ABCEDYL GA SOLUTION BUVABLE

Učinkovina:
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

Ciljne živalske vrste:
govedo
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish

Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Czech Danish Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English Italian Latvian Lithuanian Hungarian Romanian Finnish Swedish Icelandic

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Pot uporabe:
Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
139.90 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
139.90 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
139.90 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
139.90 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
139.90 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
139.90 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
139.90 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Peroralna raztopina

Karenca glede na pot uporabe:
Peroralna uporaba:

- **govedo**
  - All relevant tissues. 0 day
- **Pig**
  - All relevant tissues. 0 day
- **Rabbit**
  - All relevant tissues. 0 day
- **Horse**
  - All relevant tissues. 0 day
- **Sheep**
  - All relevant tissues. 0 day
- **Goat**
  - All relevant tissues. 0 day
- **Poultry**
  - All relevant tissues. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QV03AX

Pravni status za dobavo/izdajo zdravila:

Zdravilo, ki se izdaja brez veterinarskega recepta

Status dovoljenja za promet z zdravilom:
Valid

Ima dovoljenje za promet v:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Na voljo v:
France

Opis ovojnine zdravila:
Na voljo samo v French
Na voljo samo v French

Dodatne informacije

Vrsta dovoljenja:
Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Boiron

Datum dovoljenja za promet z zdravilom:
28/06/2012

Proizvodna mesta, odgovorna za sproščanje serij:
Boiron

Pristojni organ:
French Agency For Food, Environmental And Occupational Health & Safety

Številka dovoljenja za promet z zdravilom:
FR/V/2653261 5/2012

Datum spremembe statusa dovoljenja za promet:
21/06/2017

Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Povzetek glavnih značilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

Package Leaflet and Labelling

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.