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Broncho-Akut ReVet RV 3A - Globuli für Tiere

Ima dovoljenje za promet

- Aconitum napellus C9
- FERRUM PHOSPHORICUM C9
- HEPAR SULFURIS C9

Informacije o zdravilu

Ime zdravila:
Broncho-Akut ReVet RV 3A - Globuli für Tiere

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

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

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Pot uporabe:

Peroralna uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
3.33 milligram(s) / 1.00 gram(s)

Na voljo samo v English
3.33 milligram(s) / 1.00 gram(s)

Na voljo samo v English
3.33 milligram(s) / 1.00 gram(s)

Farmacevtska oblika:

Kroglice

Karenca glede na pot uporabe:

Peroralna uporaba:

- **Pigeon**
  - Meat and offal. 0 day
- **Turkey (hen)**
  - Eggs. 0 day
  - Meat and offal. 0 day
- **govedo**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Chicken**
  - Eggs. 0 day
  - Meat and offal. 0 day
- **Duck**

- Eggs. 0 day
- Meat and offal. 0 day

•
Goose
- Eggs. 0 day
- Meat and offal. 0 day

•
Goat
- Meat and offal. 0 day
- Milk. 0 hour

•
Sheep
- Meat and offal. 0 day
- Milk. 0 hour

•
Horse
- Meat and offal. 0 day
- Milk. 0 hour

•
Rabbit
- Meat and offal. 0 day

•
Pig
- Meat and offal. 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

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

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:
Pharmazeutische Fabrik Dr. Reckeweg & Co. GmbH

Datum dovoljenja za promet z zdravilom:
6/04/2010

Proizvodna mesta, odgovorna za sproščanje serij:
Pharmazeutische Fabrik Dr. Reckeweg & Co. GmbH

Pristojni organ:
Austrian Agency For Health And Food Safety

Številka dovoljenja za promet z zdravilom:
8-30097

Datum spremembe statusa dovoljenja za promet:
6/04/2010

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.

Označevanje

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

Navodilo za uporabo

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

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