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Pana Veyxal Salbe masť

Ima dovoljenje za promet

- Retinol palmitate
- alfa-Tocopheryl acetate
- Chymotrypsin
- Trypsin
- Papain

Informacije o zdravilu

Ime zdravila:
Pana Veyxal Salbe masť

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

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 German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

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Pot uporabe:
Dermalna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
4250.00 international unit(s) / 1.00 gram(s)

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

Na voljo samo v English
52.00 FIP / 1.00 gram(s)

Na voljo samo v English
26.00 FIP / 1.00 gram(s)

Na voljo samo v English
2.80 FIP / 1.00 gram(s)

Farmacevtska oblika:
Mazilo

Karenca glede na pot uporabe:
Dermalna uporaba:

-

govedo

- Milk. 0 day Zero days
- Meat and offal. 0 day Zero days

•

Pig

- Meat and offal. 0 day Zero days

•

Sheep

- Milk. 0 day Zero days
- Meat and offal. 0 day Zero days

•

Goat

- Milk. 0 day Zero days
- Meat and offal. 0 day Zero days

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

QD11AX

Pravni status za dobavo/izdajo zdravila:

Zdravilo, ki se izdaja na veterinarski recept

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

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Opis ovojnine zdravila:

Na voljo samo v Slovak

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 French Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Veyx Pharma GmbH

Datum dovoljenja za promet z zdravilom:
13/06/1994

Proizvodna mesta, odgovorna za sproščanje serij:
Veyx Pharma GmbH

Pristojni organ:
Institute For State Control Of Veterinary Biologicals And Medicaments

Številka dovoljenja za promet z zdravilom:
96/158/94-S

Datum spremembe statusa dovoljenja za promet:
13/06/1994

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

Dokumenti

Combined File of all Documents

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