File downloaded on 2026-10-31
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RESPIVAC aMPV (--) - Lyophilisate for oculonasal suspension/use in drinking water

Ima dovoljenje za promet

- Turkey rhinotracheitis virus, strain 1062, Live

Informacije o zdravilu

Ime zdravila:
RESPIVAC aMPV (--) - Lyophilisate for oculonasal suspension/use in drinking water

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
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

Pot uporabe:
Okulonazalno dajanje
Dajanje v vodo za pitje

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
Presentation_strength:$10^{2,4}$ – $10^{5,4}$ $CCID_{50}$/dose Reference:Ph. Eur. Index:0

Farmacevtska oblika:
Liofilizat za okulonazalno suspenzijo/dajanje v vodo za pitje

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Karenca glede na pot uporabe:
Okulonazalno dajanje:

- **Chicken**
  - Not applicable. 0 day Zero days
- **Turkey**
  - Not applicable. 0 day Zero days

Dajanje v vodo za pitje:

- **Chicken**
  - Not applicable. 0 day Zero days
- **Turkey**
  - Not applicable. 0 day Zero days

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI01AD01

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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Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

Opis ovojnine zdravila:

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

Na voljo samo v English

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 Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:

Laboratorios Hipra S.A.

Datum dovoljenja za promet z zdravilom:

30/05/2024

Proizvodna mesta, odgovorna za sproščanje serij:

Laboratorios Hipra S.A.

Pristojni organ:

European Commission

Številka dovoljenja za promet z zdravilom:
Ta podatek za to zdravilo ni na voljo.

Datum spremembe statusa dovoljenja za promet:
30/05/2024

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

Dokumenti

Combined File of all Documents

slovenščina (PDF)
Objavljeno na dan: 9/10/2026
Prenesi

ema-puar-v6160-respivacampv-vra0000309778-en.pdf

ema-puar-v6160-respivacampv-former-respivactrt-en.pdf