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Respiporc FLUpan H1N1 (--) - Suspension for injection

Ima dovoljenje za promet

- Influenza A virus, subtype H1N1, strain A/Jena/VI5258/2009 pdm09, Inactivated

Informacije o zdravilu

Ime zdravila:
Respiporc FLUpan H1N1 (--) - Suspension for injection

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

Pot uporabe:
intramuskularna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
Presentation_strength:≥ 16 HU Reference:HSE Index:0

Farmacevtska oblika:

Suspenzija za injiciranje

Karenca glede na pot uporabe:

intramuskularna uporaba:

- **Pig**

  - Not applicable. 0 day Zero days

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

QI09AA03

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

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Na voljo v:
Austria , Belgium , Bulgaria , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

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

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 Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Ceva Sante Animale

Datum dovoljenja za promet z zdravilom:
17/05/2017

Proizvodna mesta, odgovorna za sproščanje serij:
IDT Biologika GmbH
Ceva-Phylaxia Zrt.

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
17/05/2017

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

Dokumenti

ema-puar-respiporc-flupan-h1n1-v-003993-var-ii-0013-en.pdf

ema-puar-respiporc-flupan-h1n1-v-3993-par-en.pdf