

Rispoval 3 BRSV Pi3 BVD

Lyophilisate and suspension for suspension for injection for cattle

Ima
dovoljenje za
promet

- Bovine parainfluenza virus 3, strain RLB103, Live
- Bovine viral diarrhoea virus 1, strain 5960, Inactivated
- Bovine viral diarrhoea virus 1, strain 6309, Inactivated
- Bovine respiratory syncytial virus, strain 375, Live

Informacije o zdravilu

Ime zdravila:

RISPOVAL 3 BRSV PI3 BVD LYOPHILISATE AND SUSPENSION FOR SUSPENSION FOR INJECTION FOR CATTLE

Rispoval 3 BRSV Pi3 BVD Lyophilisate and suspension for suspension for injection for cattle

Učinkovina:

Na voljo samo v [English](#)

Na voljo samo v [English](#)

Na voljo samo v [English](#)

Na voljo samo v [English](#)

Ciljne živalske vrste:

govedo

Pot uporabe:

intramuskularna uporaba

Podatki o zdravilu**Učinkovina / Jakost:**

Na voljo samo v [English](#)

100000.00 50% cell culture infectious dose / 1.00 Dose

Na voljo samo v [English](#)

3.00 log2 serum neutralising unit(s) / 1.00 Dose

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Na voljo samo v [English](#)

100000.00 50% cell culture infectious dose / 1.00 Dose

Farmacevtska oblika:

Liofilizat in suspenzija za suspenzijo za injiciranje

Karenca glede na pot uporabe:**intramuskularna uporaba:**

-

govedo

- All relevant tissues. 0 day

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

QI02AH

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:

Ireland

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](#) [Lithuanian](#) [Norwegian](#)

Imetnik dovoljenja za promet z zdravilom:

Zoetis Belgium S.A.

Datum dovoljenja za promet z zdravilom:

9/12/2013

Proizvodna mesta, odgovorna za sproščanje serij:

Zoetis Belgium

Pristojni organ:

Health Products Regulatory Authority

Številka dovoljenja za promet z zdravilom:

VPA10387/060/001

Datum spremembe statusa dovoljenja za promet:

9/12/2013

Referenčna država članica:

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Številka postopka:

Zadevne države članice:

Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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[Norwegian](#)

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