

Tulaject 100 mg/ml solution for injection for cattle, pigs and sheep

Heimilað

- Tulathromycin

Auðkenni lyfs

Heiti lyfs:

Tulaject 100 mg/ml solution for injection for cattle, pigs and sheep

Virkt efni:

Aðeins fáanlegt í [enska](#)

Marktegund:

Nautgripir

Svín

Sauðkind

Leið stjórnsýslu:

Til notkunar undir húð

Til notkunar í vöðva

Upplýsingar um lyf

Virkt efni og styrkur:

Aðeins fáanlegt í [enska](#)

100.00 milligram(s) / 1.00 millilitre(s)

Lyfjaform:

Stungulyf, lausn

Afurðanýtingafrestur eftir íkomuleið:**Til notkunar undir húð:**

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Nautgripir

- Kjöt og innmatur. 22 dagar
- Mjólk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption. Do not use in pregnant animals, which are intended to produce milk for human consumption, within 2 months of expected parturition

- Kjöt og innmatur. 22 dagar
- Mjólk. no withdrawal period

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Til notkunar í vöðva:

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Svín

- Kjöt og innmatur. 13 dagar

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Sauðkind

- Kjöt og innmatur. 16 dagar
- Mjólk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption. Do not use in pregnant animals, which are intended to produce milk for human

consumption, within 2 months of expected parturition

- Kjöt og innmatur. 16 dagar
- Mjólk. no withdrawal period

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ATC flokkun (dýrallyf) (e. Anatomical therapeutic chemical veterinary - ATC-VET):

QJ01FA94

Lögformleg staða:

Ávísunarskylt dýrallyf

Staða leyfis:

Gilt

Heimilað í:

Ítalía

Lýsing umbúða:

Aðeins fáanlegt í enska

Aðeins fáanlegt í enska

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Aðrar upplýsingar

Réttindategund:

Lagalegur grundvöllur markaðsleyfis:

Aðeins fáanlegt í enska ítalska lettneska Norwegian

Markaðsleyfishafi:

Alivira Animal Health Limited

Dagsetning markaðsleyfis:

28/12/2020

Framleiðslustaður fyrir losun lotu:

Laboratorios Karizoo S.A.

Ábyrgt yfirvald:

Ministry Of Health

Markaðsleyfisnúmer:

105465

Dagsetning á breytingu stöðu:

28/12/2020

Umsjónarland (RMS):

Spánn

Ferilsnúmer:

ES/V/0373/001

Þátttökulönd (CMS):

Belgía Kýpur Eistland Frakkland Þýskaland Grikkland Ungverjaland Ítalía
Lettland Litáen Holland Pólland Portúgal Rúmenía

Til að tilkynna upplýsingar um aukaverkanir vegna dýralyfja vinsamlegast farið á
www.adrreports.eu/vet

Skjöl

Combined File of all Documents

Þetta skjal er ekki til á þessu tungumáli (íslenska). Þú getur fundið það á öðru tungumáli hér að neðan.

Samantekt á eiginleikum lyfs

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Áletranir

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