

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

Zugelassen

- Tulathromycin

Produktidentifikation

Arzneimittel:

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

Wirkstoff:

Verfügbar nur in englisch

Zieltierarten:

Rind

Schwein

Schaf

Art der Anwendung:

subkutane Anwendung

intramuskuläre Anwendung

Produktdetails

Wirkstoff und Stärke:

Verfügbar nur in englisch

100.00 milligram(s) / 1.00 millilitre(s)

Darreichungsform:

Injektionslösung

Wartezeit(en) per Anwendungsart:

subkutane Anwendung:

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Rind

- Fleisch und Innereien. 22 Tag
- Milch. 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

- Fleisch und Innereien. 22 Tag
- Milch. no withdrawal period

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intramuskuläre Anwendung:

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Schwein

- Fleisch und Innereien. 13 Tag

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Schaf

- Fleisch und Innereien. 16 Tag
- Milch. 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

- Fleisch und Innereien. 16 Tag
- Milch. no withdrawal period

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Anatomisch-therapeutisch-chemischer Veterinär-code (ATCvet-Code):

QJ01FA94

Abgaberegulierung:

Verfügbar nur in tschechisch estnisch englisch französisch italienisch lettisch litauisch portugiesisch rumänisch slowenisch finnisch schwedisch isländisch Norwegian

Zulassungsstatus:

Zulassung gültig

Zugelassen in:

Italien

Packungsbeschreibung:

Verfügbar nur in englisch

Verfügbar nur in englisch

Verfügbar nur in englisch

Zusätzliche Informationen

Anspruchstyp:

Verfügbar nur in englisch französisch kroatisch italienisch lettisch finnisch schwedisch isländisch Norwegian

Rechtsgrundlage der Produktzulassung:

Verfügbar nur in [englisch](#) [italienisch](#) [lettisch](#) [Norwegian](#)

Zulassungsinhaber:

Alivira Animal Health Limited

Zulassungsdatum:

28/12/2020

Für die Chargenfreigabe zuständige Produktionsstätten:

Laboratorios Karizoo S.A.

Zuständige Behörde:

Ministry Of Health

Zulassungsnummer:

105465

Tag der Änderung des Zulassungsstatus:

28/12/2020

Referenzmitgliedstaat:

Spanien

Verfahrensnummer:

ES/V/0373/001

Betroffene Mitgliedstaaten:

Belgien Zypern Estland Frankreich Deutschland Griechenland Ungarn Italien
Lettland Litauen Niederlande Polen Portugal Rumänien

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Dokumente

Kombinierte Datei aller Dokumente

Dieses Dokument existiert in dieser Sprache nicht (deutsch). Sie finden es in einer anderen Sprache unten.