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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:

- **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:

- **Schwein**
  - Fleisch und Innereien. 13 Tag
- **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

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

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

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

- **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

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

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

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Anatomisch-therapeutisch-chemischer Veterinärcode (ATCvet-Code):
QJ01FA94

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

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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

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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 Rumaenien

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.

Zusammenfassung der Merkmale des Arzneimittels

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

Etikettierung

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

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