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

Godkänd

- Tulathromycin

Produktens identitetsbeteckning

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

Aktiv substans:
Finns tillgänglig endast på engelska

Djurslag:
Nöt
Svin
Får

Administreringsväg:
Subkutan användning
Intramuskulär användning

Ytterligare information om produkten

Aktiv substans och styrka:
Finns tillgänglig endast på engelska
100.00 milligram(s) / 1.00 millilitre(s)

Läkemedelsform:
Injektionsvätska, lösning

Karenstid per administreringsväg:
Subkutan användning:

- **Nöt**
  - Meat and offal. 22 dygn
  - Milk. 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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Intramuskulär användning:

- **Svin**
  - Meat and offal. 13 dygn
- **Får**
  - Meat and offal. 16 dygn
  - Milk. 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

- Meat and offal. 16 dygn
- Milk. no withdrawal period

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- **Får**
  - Meat and offal. 16 dygn
  - Milk. no withdrawal period

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Anatomisk terapeutisk kemisk veterinärkod (ATCvet-kod):
QJ01FA94

Receptstatus:
Receptbelagt veterinärmedicinskt läkemedel

Godkännandestatus:
Godkänd

Godkänd i:
Italien

Förpackningsbeskrivning:
Finns tillgänglig endast på engelska
Finns tillgänglig endast på engelska
Finns tillgänglig endast på engelska

Ytterligare information

Typ av berättigande:

Legal grund för produktgodkännande:
Finns tillgänglig endast på engelska italienska lettiska Norwegian

Innehavare av godkännande för försäljning:
Alivira Animal Health Limited

Godkännandedatum:
28/12/2020

Tillverkningsplatser för frisläppande av tillverkningssatser:
Laboratorios Karizoo S.A.

Ansvarig myndighet:
Ministry Of Health

Godkännandenummer:
105465

Datum för ändring av godkännandestatus:
28/12/2020

Referensmedlemsstat:
Spanien

Procedurnummer:
ES/V/0373/001

Berörda medlemsstater:
Belgien Cypern Estland Frankrike Tyskland Grekland Ungern Italien Lettland Litauen Nederländerna Polen Portugal Rumänien

Rapporter om misstänkta biverkningar: www.adrreports.eu/vet

Dokumenter

Gemensam fil för alla dokument

Det här dokumentet finns inte på detta språk (svenska). Du kan hitta det på ett annat språk nedan.

Produktresumé

Det här dokumentet finns inte på detta språk (svenska). Du kan hitta det på ett annat språk nedan.

Märkningstext

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