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

Toegelaten

- Tulathromycin

Productidentificatie

Naam van het geneesmiddel:
Tulaject 100 mg/ml solution for injection for cattle, pigs and sheep

Werkzame stof:
Alleen beschikbaar in Engels

Doeldiersoort(en):
Rund
Varken
Schaap

Toedieningsweg:
Subcutaan gebruik
Intramusculair gebruik

Productgegevens

Werkzame stof / Sterkte:
Alleen beschikbaar in Engels

100.00 milligram(s) / 1.00 millilitre(s)

Farmaceutische vorm:
Oplossing voor injectie

Wachttijd per toedieningsweg:
Subcutaan gebruik:

- **Rund**
  - Meat and offal. 22 day
  - 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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Intramusculair gebruik:

- **Varken**
  - Meat and offal. 13 day
- **Schaap**
  - Meat and offal. 16 day
  - 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 day
- Milk. no withdrawal period

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- Meat and offal. 16 day
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- **Schaap**
  - Meat and offal. 16 day
  - 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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Anatomisch therapeutisch chemische veterinaire classificatie (ATCvet code) :
QJ01FA94

Afleverstatus:
Alleen beschikbaar in Tsjechisch Estlands Engels Frans Italiaans Letlands Litouws Portugees Roemeense Toedieningsweg Fins Zweeds IJslands Norwegian

Vergunningsstatus:
Valid

Toegelaten in:
Italië

Verpakkingsomschrijving:
Alleen beschikbaar in Engels
Alleen beschikbaar in Engels
Alleen beschikbaar in Engels

Aanvullende informatie

Type vergunning voor het in de handel:
Alleen beschikbaar in Engels Frans Kroatisch Italiaans Letlands Fins Zweeds IJslands Norwegian

Wettelijke basis productvergunning:
Alleen beschikbaar in Engels Italiaans Letlands Norwegian

Handelsvergunninghouder:
Alivira Animal Health Limited

Handelsvergunningsdatum:
28/12/2020

Locaties fabrikanten vrijgifte:
Laboratorios Karizoo S.A.

Verantwoordelijke instantie:
Ministry Of Health

Vergunningsnummer:
105465

Datum toelatingswijziging:
28/12/2020

Rapporterende lidstaat:
Spanje

Procedurenummer:
ES/V/0373/001

Betrokken lidstaten:
België Cyprus Estland Frankrijk Duitsland Griekenland Hongarije Italië Letland Litouwen Nederland Polen Portugal Roemenië

Om bijwerkingen van diergeneesmiddelen te raadplegen, zie www.adrreports.eu/vet

Documenten

Combined File of all Documents

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Samenvatting van de productkenmerken

Dit document bestaat niet in deze taal (Nederlands). U kunt het hieronder vinden in een andere taal.

Etikettering

Dit document bestaat niet in deze taal (Nederlands). U kunt het hieronder vinden in een andere taal.

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