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

Autorizado

- Tulathromycin

Identificación del medicamento

Nombre del medicamento:
Tulaject 100 mg/ml solution for injection for cattle, pigs and sheep

Principio activo:
Disponible únicamente en English

Especies de destino:
Bovino
Porcino
Ovino

Vía de administración:
Vía subcutánea
Vía intramuscular

Datos del medicamento

Principio activo y concentración:
Disponible únicamente en English

100.00 Miligramo(s) / 1.00 Mililitro(s)

Forma farmacéutica:
Solución inyectable

Tiempo de espera por vía de administración:
Vía subcutánea:

- **Bovino**
  - Meat and offal. 22 Día
  - 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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Vía intramuscular:

- **Porcino**
  - Meat and offal. 13 Día
- **Ovino**
  - Meat and offal. 16 Día
  - 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 Día
- Milk. no withdrawal period

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- Meat and offal. 16 Día
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- **Ovino**
  - Meat and offal. 16 Día
  - 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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código químico anatómico-terapéutico para medicamentos veterinarios (ATCvet):
QJ01FA94

Condiciones de dispensación:
Disponible únicamente en Czech Estonian English French Italian Latvian Lithuanian Portuguese Romanian Slovenian Finnish Swedish Icelandic Norwegian

Estado de la autorización:
Autorizado

Autorizado en:
Italia

Descripción del formato:
Disponible únicamente en English
Disponible únicamente en English
Disponible únicamente en English

Información adicional

Tipo legal de la autorización:
Disponible únicamente en English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Base legal de la autorización del medicamento:
Disponible únicamente en English Italian Latvian Norwegian

Titular de la autorización de comercialización:
Alivira Animal Health Limited

Fecha de autorización de comercialización:
28/12/2020

Fabricante responsable de la liberación del lote:
Laboratorios Karizoo S.A.

Autoridad responsable:
Ministry Of Health

Número de autorización:
105465

Fecha de modificación del estado de la autorización:
28/12/2020

Estado miembro de referencia:
España

Número de procedimiento:
ES/V/0373/001

Estados miembros afectados:
Bélgica Chipre Estonia Francia Alemania Grecia Hungría Italia Letonia Lituania Países Bajos Polonia Portugal Rumania; Rumanía

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Documentos

Combined File of all Documents

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Ficha técnica o resumen de las características del producto

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Etiquetado

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