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

Autorisé

- Tulathromycin

Identification du produit

Dénomination du médicament:
Tulaject 100 mg/ml solution for injection for cattle, pigs and sheep

Substance(s) active(s):
Disponible uniquement en Anglais

Espèces cibles:
Bovins
Porc
Mouton

Voie d’administration:
Voie sous-cutanée
Voie intramusculaire

Informations sur le produit

Substance(s) active(s) / Concentration:
Disponible uniquement en Anglais
100.00 milligram(s) / 1.00 millilitre(s)

Forme pharmaceutique:
Solution injectable

Temps d'attente par voie d'administration:
Voie sous-cutanée:

- **Bovins**
  - Viande et abats. 22 day
  - Lait. 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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Voie intramusculaire:

- **Porc**
  - Viande et abats. 13 day
- **Mouton**
  - Viande et abats. 16 day
  - Lait. 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

- Viande et abats. 16 day
- Lait. no withdrawal period

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- **Mouton**
  - Viande et abats. 16 day
  - Lait. no withdrawal period

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Code anatomique, thérapeutique, chimique, vétérinaire (ATCvet):
QJ01FA94

Conditions de délivrance:
Médicament vétérinaire soumis à prescription vétérinaire

Statut de l’autorisation:
Valide

Autorisé en:
Italie

Description des conditionnements:
Disponible uniquement en Anglais
Disponible uniquement en Anglais
Disponible uniquement en Anglais

Informations complémentaires

Type d'autorisation:

Base légale de l'autorisation du produit:
Disponible uniquement en Anglais Italian Latvian Norwegian

Titulaire de l’autorisation de mise sur le marché:
Alivira Animal Health Limited

Date de l'autorisation de mise sur le marché:
28/12/2020

Site(s) de libération des lots du produit fini:
Laboratorios Karizoo S.A.

Autorité responsable:
Ministry Of Health

Numéro de l’autorisation:
105465

Date de modification du statut de l’autorisation:
28/12/2020

État membre de référence:
Espagne

Numéro de procédure:
ES/V/0373/001

États membres concernés:
Belgique Chypre Estonie France Allemagne Grèce Hongrie Italie Lettonie Lituanie Pays-Bas Pologne Portugal Roumanie

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Documents

Fichier combinant tous les documents

Ce document n'existe pas dans cette langue (français). Vous pouvez le trouver dans une autre langue ci-dessous.

Résumé des caractéristiques du produit

Ce document n'existe pas dans cette langue (français). Vous pouvez le trouver dans une autre langue ci-dessous.

Etiquetage

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