

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:**

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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:

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Porc

- Viande et abats. 13 day

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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

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- Viande et abats. 16 day
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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:

Espagne

Description des conditionnements:

Disponible uniquement en [Spanish](#)

Disponible uniquement en [Spanish](#)

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Informations complémentaires

Type d'autorisation:

Marketing Authorisation

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:

4/12/2020

Site(s) de libération des lots du produit fini:

Laboratorios Karizoo S.A.

Autorité responsable:

Spanish Agency Of Medicines And Medical Devices

Numéro de l'autorisation:

3950 ESP

Date de modification du statut de l'autorisation:

5/12/2020

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

Pour connaître les évènements indésirables rapportés concernant les médicaments vétérinaires, consulter ce site www.adrreports.eu/vet

Documents

Résumé des caractéristiques du produit

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

Notice

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

Etiquetage

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

Fichier combinant tous les documents

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