

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

Autorisert

- Tulathromycin

Produkt identifikasjon

Legemidlets navn:

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

Aktiv substans virkestoff:

Bare tilgjengelig i [Engelsk](#)

Målarter:

storfe

gris

sau

Administrasjonsvei:

Subkutan bruk

Intramuskulær bruk

Produktdetaljer

Aktiv substans og styrke:

Bare tilgjengelig i [Engelsk](#)

100.00 milligram / 1.00 milliliter

Legemiddelform:

Injeksjonsvæske, oppløsning

Tilbakeholdelsestid etter administrasjonsvei:**Subkutan bruk:**

-

storfe

- Slakt. 22 dag
- Melk. 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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Intramuskulær bruk:

-

gris

- Slakt. 13 dag

-

sau

- Slakt. 16 dag
- Melk. no withdrawal period

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sau

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Anatomisk terapeutisk kjemisk klassifisering for veterinærpreparater (ATCvet):

QJ01FA94

Juridisk status for forsyning:

Veterinært legemiddel gjenstand for veterinær forskrivning

Status for markedsføringstillatelse:

Gyldig

Autorisert i:

IT

Pakningsbeskrivelse:

Bare tilgjengelig i [Engelsk](#)

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Tilleggsinformasjon

Rettighetstype:

Juridisk grunnlag for markedsføringstillatelse:

Generisk søknad (Artikkel 13(1) i Direktiv 2001/82/EC)

Innehaver av markedsføringstillatelse:

Alivira Animal Health Limited

Markedsføringsgodkjenningsdato:

28/12/2020

Tilvirker for batchfrigivelse:

Laboratorios Karizoo S.A.

Ansvarlig myndighet:

Ministry Of Health

Godkjenningsnummer:

105465

Status for endring av markedsføringstillatelse:

28/12/2020

Referanse medlemsstat:

ES

Prosedyrenummer:

ES/V/0373/001

Gjeldende medlemsstater:

BE Kypros EE FR DE EL HU IT LV LT NL PL PT RO

Rapporter om mistenkte bivirkninger: www.adrreports.eu/vet

Dokumenter

Combined File of all Documents

Dette dokumentet eksisterer ikke på dette språket (Norsk). Du kan finne det på et annet språk nedenfor.

Summary of Product Characteristics

Dette dokumentet eksisterer ikke på dette språket (Norsk). Du kan finne det på et annet språk nedenfor.

Labelling

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