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Flortek 300 mg/ml solution for injection for cattle, sheep and pigs

Autoriseret

- Florfenicol

Produktidentifikation

Lægemiddelnavn:
Flortek 300 mg/ml solution for injection for cattle, sheep and pigs

Aktiv substans:
Kun tilgængelig på English

Dyreart:
Kvæg
Får
Svin

Administrationsvej:
Intramuskulær anvendelse
Subkutan anvendelse

Produktoplysninger

Aktiv substans og styrke:
Kun tilgængelig på English

300.00 milligram(s) / 1.00 millilitre(s)

Lægemiddelform:
Injektionsvæske, opløsning

Tilbageholdelsestid efter administrationsrute:
Intramuskulær anvendelse:

- **Kvæg**
  - Meat and offal. no withdrawal period IM use: 30 days. SC use: 44 days.
  - Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption including pregnant animals intended to produce milk for human consumption.

- **Får**
  - Meat and offal. 39 dag
  - Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption including pregnant animals intended to produce milk for human consumption

- **Svin**
  - Meat and offal. 18 dag

- **Kvæg**
  - Meat and offal. no withdrawal period IM use: 30 days. SC use: 44 days.
  - Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption including pregnant animals intended to produce milk for human consumption.

- **Får**

- Meat and offal. 39 dag

- Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption including pregnant animals intended to produce milk for human consumption

Subkutan anvendelse:

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  - Milk. no withdrawal period

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Anatomisk terapeutisk kemisk veterinær (ATCvet) kode:
QJ01BA90

Udleveringsbestemmelse:
Kun tilgængelig på Czech Estonian English French Italian Latvian Lithuanian Portuguese Romanian Slovenian Finnish Swedish Icelandic Norwegian

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Godkendelsesstatus:
Gyldig

Autoriseret i:
Kun tilgængelig på Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Pakningsbeskrivelse:

Kun tilgængelig på English
Kun tilgængelig på English

Yderligere oplysninger

Berettigelsestype:
Kun tilgængelig på English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Lovgrundlag for godkendelse:
Kun tilgængelig på English Italian Latvian Norwegian

Indehaver af markedsføringstilladelsen:
Laboratorios Karizoo S.A.

Dato for markedsføringstilladelse:
1/01/2023

Fremstillere ansvarlige for batchfrigivelse:
Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Ansvarlig myndighed:
Ministry Of Health

Markedsføringstilladelsesnummer:
105642

Dato for ændring af godkendelsesstatus:
1/01/2023

Referencemedlemsstat:
Kun tilgængelig på Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Procedurenummer:
ES/V/0409/001

Berørte medlemsstater:

Kun tilgængelig på Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Indberetninger om formodede bivirkninger på veterinærlægemidler: www.adrreports.eu/vet

Dokumenter

Combined File of all Documents

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Produktresumé

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Indlægsseddel

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Etikettering

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