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CADOREX 300 mg/ml Solution for injection for cattle, pigs and sheep

Autoriseret

- Florfenicol

Produktidentifikation

Lægemiddelnavn:
CADOREX 300 mg/ml Solution for injection for cattle, pigs and sheep

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

Meat and offal: IM: 30 dAYS / SC: 44 Days

  - Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption.

- **Får**
  - Meat and offal. no withdrawal period

Meat and offal: im 39 Days

  - Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption.

- **Svin**
  - Meat and offal. no withdrawal period

Meat and offal: Intramuscular use: 18 days

- **Kvæg**
  - Meat and offal. no withdrawal period

Meat and offal: IM: 30 dAYS / SC: 44 Days

  - Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption.

- **Får**

- Meat and offal. no withdrawal period Meat and offal: im 39 Days

- Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption.

Subkutan anvendelse:

- **Kvæg**
  - Meat and offal. no withdrawal period

Meat and offal: IM: 30 dAYS / SC: 44 Days

- Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption.

- **Kvæg**
  - Meat and offal. no withdrawal period

Meat and offal: IM: 30 dAYS / SC: 44 Days

- Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption.

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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Tilgængelig i:

Netherlands

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:
Industrial Veterinaria S.A.

Dato for markedsføringstilladelse:
15/08/2017

Fremstillere ansvarlige for batchfrigivelse:
Industrial Veterinaria S.A.
aniMedica GmbH

Ansvarlig myndighed:
Medicines Evaluation Board

Markedsføringstilladelsesnummer:
REG NL 120675

Dato for ændring af godkendelsesstatus:
2/02/2022

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

Procedurenummer:
ES/V/0246/001

Berørte medlemsstater:

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Indberetninger om formodede bivirkninger på veterinærlægemidler: www.adrreports.eu/vet

Dokumenter

Combined File of all Documents

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Etikettering

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Indlægsseddel

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