

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

Autorisert

- Florfenicol

Product identification

Legemidlets navn:

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

FLORFENIS 300 mg/ml solutie injectabila pentru bovine, ovine si porcine

Virkestoff:

Tilgjengelig bare i [English](#)

Dyrearter:

storfe

gris

sau

Administrering:

Intramuskulær bruk

Subkutan bruk

Product details

Virkestoff / Styrke:

Tilgjengelig bare i [English](#)

300.00 milligram / 1.00 milliliter

Legemiddelform:

Withdrawal period by route of administration:

Intramuskulær bruk:

• **storfe**

- Slakt. no withdrawal period 20 mg / kg bw, twice: 30 days
- Melk. no withdrawal period

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

• **gris**

- Slakt. 18 dag

• **sau**

- Slakt. 39 dag
- Melk. no withdrawal period

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

Subkutan bruk:

• **storfe**

- Slakt. no withdrawal period 40 mg / kg bw, once: 44 days
- Melk. no withdrawal period

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

Anatomisk terapeutisk kjemisk klassifisering for veterinærpreparater (ATCvet):

QJ01BA90

Utleveringsbestemmelser :

Veterinært legemiddel gjenstand for veterinær forskrivning

Status for markedsføringstillatelse:

Gyldig

Authorised in:

RO

Pakningsvedlegg:

Tilgjengelig bare i [English](#)

Tilgjengelig bare i [English](#)

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Additional information

Entitlement type:

Marketing Authorisation

Rettslig grunnlag for produktgodkjenning:

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

Innehaver av markedsføringstillatelse:

Laboratorios Syva S.A.

Marketing authorisation date:

10/05/2021

Tilvirker ansvarlig for batchfrigivelse:

Laboratorios Syva S.A.U.

Ansvarlig myndighet:

Institute For Control Of Biological Products And Veterinary Medicines

Godkjenningsnummer:

210059

Dato for endring av status for markedsføringstillatelse:

10/05/2021

Referanse medlemsstat:

ES

Prosedyrenummer:

ES/V/0377/001

Gjeldende medlemsstater:

BE DK FR DE EL HU Irland IT LT NL PL PT RO

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

eu-PUAR-florfenis-300-mg-ml-solution-for-injection-for-cattle--sheep-and-pigs-en.pdf

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