

QUINOLCEN 100 mg/ml Injection

Authorised

- Enrofloxacin

Product identification

Medicine name:

QUINOLCEN 100 mg/ml Injection

Colmyc 100 mg/ml oldatos injekció szarvasmarhák, juhok, kecskék és sertések részére A.U.V.

Active substance:

Enrofloxacin

Target species:

Cattle

Sheep

Goat

Pig

Route of administration:

Subcutaneous use

Intravenous use

Intramuscular use

Product details

Active substance and strength:

Enrofloxacin

100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

-

Cattle

- Meat and offal. no withdrawal period

Meat and offal: IV: 5 days. Via SC: 12 days

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Sheep

- Meat and offal. 4 day

-

Goat

- Meat and offal. 6 day

-

Cattle

- Milk. no withdrawal period Milk: Via IV: 3 days. Via SC: 4 days;

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Sheep

- Milk. 3 day

-

Goat

- Milk. 4 day

Intravenous use:

-

Cattle

- Meat and offal. no withdrawal period

Meat and offal: IV: 5 days. Via SC: 12 days

-

Cattle

- Milk. no withdrawal period
- Milk: Via IV: 3 days. Via SC: 4 days;

Intramuscular use:

-

Pig

- Meat and offal. 13 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Hungary

Package description:

box containing 1 vial of 250 ml

box containing 1 vial of 100 ml

box containing 1 vial of 50 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

S P Veterinaria S.A.

Marketing authorisation date:

5/05/2010

Manufacturing sites for batch release:

S P Veterinaria S.A.

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

2728/X/10 MgSzH ÁTI

Date of authorisation status change:

5/05/2010

Reference member state:

Spain

Procedure number:

ES/V/0150/001

Concerned member states:

Belgium Bulgaria Greece Hungary Italy Luxembourg Poland Portugal
Romania

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

Documents

Combined File of all Documents