

KELACYL 100 mg/ml solution for injection for cattle and pigs

Authorised

- Marbofloxacin

Product identification

Medicine name:

KELACYL 100 mg/ml solution for injection for cattle and pigs
Kelacyl

Active substance:

Marbofloxacin

Target species:

Cattle
Pig (sow)

Route of administration:

Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

Marbofloxacin
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Withdrawal period by route of administration:

Intramuscular use:

• **Cattle**

- Meat and offal. no withdrawal period

Meat and offal: Respiratory: 3 Days (8 mg/kg (IV/SC/IM)) 6 Days (2 mg/kg (IM)) /
Mastitis 3 Days (2 mg/kg (IV/IM/SC))

- Milk. no withdrawal period

Milk: Respiratory: 72 Hours (8 mg/g (IM) and 36 Hours (2 mg/g (IV / SC / IM)) /
Mastitis: 36 Hours 2 mg/g (IV / IM / SC)

• **Pig (sow)**

- Meat and offal. 4 day

Intravenous use:

• **Cattle**

- Meat and offal. no withdrawal period

Meat and offal: Respiratory: 3 Days (8 mg/kg (IV/SC/IM)) 6 Days (2 mg/kg (IM)) /
Mastitis 3 Days (2 mg/kg (IV/IM/SC))

- Milk. no withdrawal period

Milk: Respiratory: 72 Hours (8 mg/g (IM) and 36 Hours (2 mg/g (IV / SC / IM)) /
Mastitis: 36 Hours 2 mg/g (IV / IM / SC)

• **Pig (sow)**

- Meat and offal. 4 day

Subcutaneous use:

• **Cattle**

- Meat and offal. no withdrawal period

Meat and offal: Respiratory: 3 Days (8 mg/kg (IV/SC/IM)) 6 Days (2 mg/kg (IM)) /
Mastitis 3 Days (2 mg/kg (IV/IM/SC))

- Milk. no withdrawal period

Milk: Respiratory: 72 Hours (8 mg/g (IM) and 36 Hours (2 mg/g (IV / SC / IM)) /
Mastitis: 36 Hours 2 mg/g (IV / IM / SC)

• **Pig (sow)**

- Meat and offal. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA93

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Germany

Available in:

Germany

Package description:

box containing 12 vials of 250 ml individually packed in a carton box

box containing 12 vials of 100 ml individually packed in a carton box

box containing 10 vials of 250 ml individually packed in a carton box

box containing 10 vials of 100 ml individually packed in a carton box

box containing 6 vials of 250 ml individually packed in a carton box

box containing 6 vials of 100 ml individually packed in a carton box

cardboard box containing 1 vial of 250 ml

cardboard box containing 1 vial of 100 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:

Kela Kempisch Laboratorium Kela Laboratoria

Marketing authorisation date:

28/02/2013

Manufacturing sites for batch release:

Kela - Kempisch Laboratorium - Kela Laboratoria

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

401770.00.00

Date of authorisation status change:

27/06/2018

Reference member state:

Spain

Procedure number:

ES/V/0189/001

Concerned member states:

Belgium Germany Luxembourg Poland United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Package Leaflet

Labelling

eu-PUAR-esv0189001-dcp-kelacyl-100-mg-ml-en.pdf

Source URL: <https://medicines.health.europa.eu/veterinary/600000039123>