

Lignocaine HCl 2 % + Adrenaline Tartr. 0,0036 % Kela 20 mg/ml - 0.036 mg/ml Oplossing voor injectie

Authorised

- Lidocaine hydrochloride
- EPINEPHRINE BITARTRATE

Product identification

Medicine name:

Lignocaine HCl 2 % + Adrenaline Tartr. 0,0036 % Kela 20 mg/ml - 0.036 mg/ml
Oplossing voor injectie

Active substance:

Lidocaine hydrochloride
EPINEPHRINE BITARTRATE

Target species:

Cat
Horse
Dog

Route of administration:

Epidural use
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

Lidocaine hydrochloride

20.00 milligram(s) / 1.00 millilitre(s)

EPINEPHRINE BITARTRATE

0.04 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Epidural use:

-

Horse

- Meat and offal. 3 day

Intramuscular use:

-

Horse

- Meat and offal. 3 day

Subcutaneous use:

-

Horse

- Meat and offal. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN01BB52

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Package description:

Lignocaine HCl 2 % + Adrenaline Tartr. 0,0036 % Kela 20 mg/ml - 0.036 mg/ml
Solution for injection Vial of 250 ml

Lignocaine HCl 2 % + Adrenaline Tartr. 0,0036 % Kela 20 mg/ml - 0.036 mg/ml
Solution for injection Vial of 100 ml

Lignocaine HCl 2 % + Adrenaline Tartr. 0,0036 % Kela 20 mg/ml - 0.036 mg/ml
Solution for injection Vial of 50 ml

Lignocaine HCl 2 % + Adrenaline Tartr. 0,0036 % Kela 20 mg/ml - 0.036 mg/ml
Solution for injection Vial of 30 ml

Lignocaine HCl 2 % + Adrenaline Tartr. 0,0036 % Kela 20 mg/ml - 0.036 mg/ml
Solution for injection 12 Vials of 30 ml

Lignocaine HCl 2 % + Adrenaline Tartr. 0,0036 % Kela 20 mg/ml - 0.036 mg/ml
Solution for injection 12 Vials of 50 ml

Lignocaine HCl 2 % + Adrenaline Tartr. 0,0036 % Kela 20 mg/ml - 0.036 mg/ml
Solution for injection 12 Vials of 100 ml

Lignocaine HCl 2 % + Adrenaline Tartr. 0,0036 % Kela 20 mg/ml - 0.036 mg/ml
Solution for injection 12 Vials of 250 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

KELA Kempisch Laboratorium Kela Laboratoria

Marketing authorisation date:

1/05/1979

Manufacturing sites for batch release:

KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V112734

Date of authorisation status change:

27/09/2019

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

Documents

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.