LIDOCAINE, 2%, Injekční roztok

Authorised

- Lidocaine hydrochloride
- Noradrenaline tartrate

Product identification

Medicine name:
LIDOCAINE, 2%, Injekční roztok

Active substance:
Lidocaine hydrochloride
Noradrenaline tartrate

Target species:
Cat
Dog

Route of administration:
Epidural use
Perineural use
Infiltration
Topical

Product details

Active substance and strength:
Lidocaine hydrochloride

20.00 milligram(s) / 1.00 millilitre(s)

Noradrenaline tartrate
0.02 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN01BB52

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Czechia

Package description:
Available only in Czech

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:
Fatro S.p.A.

Marketing authorisation date:
9/09/2003

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/071/03-C

Date of authorisation status change:
9/11/2009

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

Documents

Summary of Product Characteristics

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Package Leaflet

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.