Nalgosed, 10mg/ml, Solution for injection

Authorised

- Butorphanol

Product identification

Medicine name:
Nalgosed, 10mg/ml, Solution for injection
Nalgosed 10 mg/ml šķīdums injekcijām

Active substance:
Butorphanol

Target species:
Cat
Dog
Horse

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Butorphanol
10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cat**
- **Dog**

Intravenous use:

- **Horse**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Cat**
- **Dog**

Subcutaneous use:

- **Cat**
- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN02AF01

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

Authorisation status:
Valid

Authorised in:
Latvia

Package description:
Glass Vial 1 x 10.0 millilitre(s)

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:
Bioveta a.s.

Marketing authorisation date:
4/04/2018

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/MRP/18/0021

Date of authorisation status change:
4/04/2018

Reference member state:
Czechia

Procedure number:
CZ/V/0142/001

Concerned member states:
Bulgaria Estonia Greece Hungary Latvia Lithuania Poland Romania Slovakia

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.
