

# Nalgosed, 10mg/ml, Solution for injection

Authorised

- Butorphanol

## Product identification

**Medicine name:**

Nalgosed, 10mg/ml, Solution for injection

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**Active substance:**

Butorphanol

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**Target species:**

Cat

Dog

Horse

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**Route of administration:**

Intramuscular use

Intravenous use

Subcutaneous use

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## Product details

**Active substance and strength:**

Butorphanol

10.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Intravenous use:**

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**Horse**

- Milk. 0 day
  - Meat and offal. 0 day
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QN02AF01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Slovakia

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**Package description:**

Glass Vial 1 x 10.0 millilitre(s)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Bioveta a.s.

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**Marketing authorisation date:**

17/10/2019

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**Manufacturing sites for batch release:**

Bioveta a.s.

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

96/035/MR/19-S

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**Date of authorisation status change:**

17/10/2019

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**Reference member state:**

Czechia

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**Procedure number:**

CZ/V/0142/001

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**Concerned member states:**

Bulgaria Estonia Greece Hungary Latvia Lithuania Poland Romania Slovakia

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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

Summary of Product Characteristics

Package Leaflet

Labelling

eu-puar-czv0142001-mr-nalgosed-en.pdf