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Nalgosed, 10mg/ml, Solution for injection

Authorised

- Butorphanol

Product identification

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

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:

Intravenous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN02AF01

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

Authorisation status:
Valid

Authorised in:
Bulgaria

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:

10/06/2018

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2817

Date of authorisation status change:
19/05/2021

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

English (PDF)
Published on: 14/03/2026
Download

Package Leaflet and Labelling

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

Package Leaflet

Labelling

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