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:
Slovakia

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:

17/10/2019

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/035/MR/19-S

Date of authorisation status change:
17/10/2019

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

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