

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
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02AF01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

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:

31/05/2018

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

National Organization For Medicines

Authorisation number:

74783/04-08-2021/K-0232301

Date of authorisation status change:

3/08/2021

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

Package Leaflet

Labelling

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