

Nalgosed, 10mg/ml, Solution for injection

Authorised

- Butorphanol

Product identification

Medicine name:

Nalgosed, 10mg/ml, Solution for injection

NALGOSED 10 mg/ml injekčný roztok

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

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