

BUTOMIDOR, 10mg/ml, Injekční roztok

Authorised

- Butorphanol

Product identification

Medicine name:

BUTOMIDOR, 10mg/ml, Injekční roztok

Active substance:

Butorphanol

Target species:

Dog

Cat

Horse

Route of administration:

Intravenous use

Subcutaneous use

Intramuscular 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

- 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:

Czechia

Available in:

Czechia

Package description:

Available only in [Czech](#)

Available only in [Czech](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Vetviva Richter GmbH

Marketing authorisation date:

8/10/2003

Manufacturing sites for batch release:

Vetviva Richter GmbH

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/065/03-C

Date of authorisation status change:

8/10/2003

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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