

Torphadine 10 mg/ml Solution for Injection for Dogs, Cats and Horses

Authorised

- Butorphanol tartrate

Product identification

Medicine name:

Torphadine 10 mg/ml Solution for Injection for Dogs, Cats and Horses

Active substance:

Butorphanol tartrate

Target species:

Horse

Dog

Cat

Route of administration:

Intramuscular use

Intravenous use

Subcutaneous use

Product details

Active substance and strength:

Butorphanol tartrate
14.58 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Horse

- Meat and offal. no withdrawal period
Withdrawal period is 0 days

Intravenous use:

-

Horse

- Meat and offal. no withdrawal period
Withdrawal period is 0 days

Subcutaneous use:

-

Horse

- Meat and offal. no withdrawal period
Withdrawal period is 0 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02AF01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Finland

Available in:

Finland

Package description:

Clear type I glass vial of 10 ml closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box

Clear type I glass vial of 20 ml closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box

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:

Le Vet. Beheer B.V.

Marketing authorisation date:

19/07/2017

Manufacturing sites for batch release:

Produlab Pharma B.V.

Responsible authority:

Finnish Medicines Agency

Authorisation number:

33653

Date of authorisation status change:

19/07/2017

Reference member state:

Netherlands

Procedure number:

NL/V/0316/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland
France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania
Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain
Sweden United Kingdom (Northern Ireland)

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.