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
Torphadine vet 10 mg/ml šķīdums injekcijām suņiem, kaķiem un zirgiem

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:

Latvia

Package description:
Clear type I glass vial of 20 ml closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box
Clear type I glass vial of 10 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:
4/10/2016

Manufacturing sites for batch release:
Produlab Pharma B.V.

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/DCP/16/0032

Date of authorisation status change:
4/10/2016

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

Combined File of all Documents

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