

# Butorgesic 10 mg/ml solution for injection

Authorised

- Butorphanol tartrate
- Butorphanol tartrate

## Product identification

### Medicine name:

Butorgesic 10 mg/ml solution for injection  
Butorgesic vet 10 mg/ml injektioneste, liuos

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### Active substance:

Butorphanol tartrate  
Butorphanol tartrate

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### Target species:

Cat  
Dog  
Horse

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### Route of administration:

Intravenous use  
Intramuscular use  
Subcutaneous use

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## Product details

### Active substance and strength:

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

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

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:**

**Intravenous use:**

- **Cat**
- **Dog**
- **Horse**
  - Milk. 0 day
  - Milk. 0 day
  - Milk. 0 day
  - Milk. 0 day
  - Meat and offal. 0 day
  - Meat and offal. 0 day
  - Meat and offal. 0 day
  - Meat and offal. 0 day

**Intramuscular use:**

- **Dog**
- **Horse**
- **Cat**

**Subcutaneous use:**

- **Dog**
  - **Horse**
  - **Cat**
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QN02AF01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Finland

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**Available in:**

Finland

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**Package description:**

Available only in [Danish](#)

Available only in [Danish](#)

Available only in [Danish](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

CP-Pharma Handelsgesellschaft mbH

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**Marketing authorisation date:**

15/01/2021

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**Manufacturing sites for batch release:**

Cp-Pharma Handelsgesellschaft mbH

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**Responsible authority:**

Finnish Medicines Agency

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**Authorisation number:**

37613

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**Date of authorisation status change:**

15/01/2021

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**Reference member state:**

Denmark

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**Procedure number:**

DK/V/0124/001

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**Concerned member states:**

Austria Finland France Hungary Italy Norway Sweden

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Summary of Product Characteristics

English (PDF)

Published on: 25/01/2022

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### Package Leaflet

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