

# Butomidor 10 mg/ml Injektionslösung für Pferde, Hunde und Katzen

Authorised

- Butorphanol tartrate

## Product identification

### Medicine name:

Butomidor 10 mg/ml Injektionslösung für Pferde, Hunde und Katzen  
BUTADOR 10 MG/ML SOLUTION INJECTABLE POUR CHEVAUX, CHIENS ET CHATS

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### Active substance:

Butorphanol tartrate

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### Target species:

Dog  
Horse  
Cat

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### Route of administration:

Intramuscular use  
Intravenous use  
Subcutaneous use

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## Product details

### Active substance and strength:

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:**

**Intramuscular use:**

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**Dog**

**Intravenous use:**

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**Dog**

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**Horse**

- Meat and offal. 0 day

- Milk. 0 hour

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**Cat**

**Subcutaneous use:**

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**Dog**

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**Cat**

- Meat and offal. 0 day

- Milk. 0 hour

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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:**

France

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**Available in:**

France

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**Package description:**

Available only in [German](#)

Available only in [German](#)

Available only in [German](#)

Available only in [German](#)

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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:**

Vetviva Richter GmbH

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**Marketing authorisation date:**

21/08/2017

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**Manufacturing sites for batch release:**

Vetviva Richter GmbH

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/1461733 9/2017

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**Date of authorisation status change:**

30/06/2022

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**Reference member state:**

Austria

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**Procedure number:**

AT/V/0005/001

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**Concerned member states:**

Belgium Denmark Finland France Germany Greece Iceland Ireland Latvia  
Lithuania Netherlands Norway Portugal Slovenia Spain Sweden  
United Kingdom (Northern Ireland)

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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

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

Package Leaflet and Labelling

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

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