

# Bupredine Multidose 0.3 mg/ml Solution for Injection for Dogs, Cats and Horses

Authorised

- Buprenorphine hydrochloride

## Product identification

**Medicine name:**

Bupredine Multidose 0.3 mg/ml Solution for Injection for Dogs, Cats and Horses  
Bupredine Multidose 0.3 mg/ml Solution for Injection for Dogs, Cats and Horses

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**Active substance:**

Buprenorphine hydrochloride

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**Target species:**

Dog  
Horse  
Cat

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**Route of administration:**

Intramuscular use  
Intravenous use

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## Product details

**Active substance and strength:**

Buprenorphine hydrochloride  
0.32 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QN02AE01

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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:**

United Kingdom (Northern Ireland)

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**Package description:**

Clear type I glass vials closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box. Pack size: 20 ml.

Clear type I glass vials closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box. Pack size: 100 ml

Clear type I glass vials closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box. Pack size: 10 ml.

Clear type I glass vials closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box. Pack size: 50 ml

Clear type I glass vials closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box. Pack size: 5 ml.

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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:**

Le Vet. B.V.

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**Marketing authorisation date:**

4/11/2015

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**Manufacturing sites for batch release:**

Produlab Pharma B.V.

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**Responsible authority:**

The Veterinary Medicines Directorate

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**Authorisation number:**

Vm 41821/3019

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**Date of authorisation status change:**

23/03/2022

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0314/001

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**Concerned member states:**

Austria Belgium Croatia Cyprus Czechia Denmark Estonia France Germany  
Greece Hungary Iceland Latvia Lithuania Luxembourg Norway Poland  
Portugal Romania Slovakia 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)