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# 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

**Active substance:**
Buprenorphine hydrochloride

**Target species:**
Dog
Cat
Horse

**Route of administration:**
Intramuscular use
Intravenous use

## Product details

**Active substance and strength:**
Buprenorphine hydrochloride

0.32 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**
Solution for injection

**Withdrawal period by route of administration:**
**Intravenous use:**

- **Horse**
  - All relevant tissues. no withdrawal period

Not authorised for use in horses intended for human consumption.

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QN02AE01

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Hungary

**Package description:**
Clear type I glass vials closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box. Pack size: 5 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: 10 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: 20 ml.

## 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:**
1/12/2015

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

**Responsible authority:**
Directorate Of Veterinary Medicinal Products

**Authorisation number:**
3702/X/15 NÉBIH ÁTI

**Date of authorisation status change:**
1/12/2015

**Reference member state:**
Netherlands

**Procedure number:**
NL/V/0314/001

**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)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

## Documents

Combined File of all Documents