

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

Horse

Cat

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02AE01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Denmark

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:

11/11/2015

Manufacturing sites for batch release:

Produlab Pharma B.V.

Responsible authority:

Danish Medicines Agency

Authorisation number:

54940

Date of authorisation status change:

11/11/2015

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

Summary of Product Characteristics

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