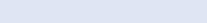
Buprecare Multidose 0.3 mg/ml Solution for Injection for Dogs and Cats

Authorised



Buprenorphine hydrochloride

Product identification

Medicine name:

Buprecare Multidose 0.3 mg/ml Solution for Injection for Dogs and Cats Buprecare Multidose 0,3 mg/ml raztopina za injiciranje za pse in mačke

Active substance:

Buprenorphine hydrochloride

Target species:

Dog 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

Withdrawal period by route of administration:

Intramuscular use:

- . Dog
- . Cat

Intravenous use:

- . Dog
- Cat

Anatomical therapeutic chemical veterinary (ATCvet) codes: ON02AE01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovenia

Package description:

Presented in a 10 ml amber Type I glass vial with a bromobutyl rubber stopper and flip-off aluminium cap.Pack size: 1 vial with 10 ml solution for injection.

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:

Ecuphar

Marketing authorisation date:

10/05/2018

Manufacturing sites for batch release:

Produlab Pharma B.V.

Responsible authority:

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number: MR/V/0622/001

Date of authorisation status change:

10/05/2018

Reference member state:

Ireland

Procedure number:

IE/V/0453/002

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus France Germany Greece Hungary Luxembourg Netherlands Poland Portugal Romania Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF) Published on: 3/05/2024 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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