

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 oldatos injekció kutyáknak és macsḱaknak A.U.V.

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

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:

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:

14/12/2017

Manufacturing sites for batch release:

Produlab Pharma B.V.

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

3914/X/17 NÉBIH ÁTI

Date of authorisation status change:

14/12/2017

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

Combined File of all Documents