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

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:
Ireland

Available in:
Ireland

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:
16/12/2011

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

Responsible authority:

Health Products Regulatory Authority

Authorisation number:
VPA10491/004/002

Date of authorisation status change:
16/12/2011

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