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Buprefelican Multidose 0.3 mg/ml solution for injection for dogs and cats

Authorised

- Buprenorphine hydrochloride

Product identification

Medicine name:
Buprefelican 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:
Finland

Available in:
Finland

Package description:
Clear type I glass vial closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box.Pack size: 50 ml.
Clear type I glass vial closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box.Pack size: 20 ml.
Clear type I glass vial closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box.Pack size: 10 ml.
Clear type I glass vial closed with a coated bromobutyl rubber stopper and aluminium cap in a carton box.Pack size: 5 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:
18/02/2016

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

Responsible authority:
Finnish Medicines Agency

Authorisation number:
32748

Date of authorisation status change:
18/02/2016

Reference member state:
Ireland

Procedure number:
IE/V/0623/001

Concerned member states:
Finland Italy 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