Bupaq Multidose 0,3 mg/ml Injektionslösung für Hunde und Katzen

Authorised

- Buprenorphine hydrochloride

Product identification

Medicine name:
Bupaq Multidose 0,3 mg/ml Injektionslösung für Hunde und Katzen

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:
Denmark

Available in:
Denmark

Package description:
Amber glass vials type I, bromobutyl rubber stopper type I, coated, aluminium cap
Package size: 10 ml
Amber glass vials type I, bromobutyl rubber stopper type I, coated, aluminium cap
Package size: 10 x 10 ml
Amber glass vials type I, bromobutyl rubber stopper type I, coated, aluminium cap
Package size: 5 x 10 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:
Vetviva Richter GmbH

Marketing authorisation date:
11/12/2013

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Danish Medicines Agency

Authorisation number:
52982

Date of authorisation status change:
11/12/2013

Reference member state:
Austria

Procedure number:
AT/V/0008/001

Concerned member states:
Belgium Bulgaria Czechia Denmark Estonia Finland France Germany Hungary Ireland Italy Latvia Lithuania Netherlands Norway Poland Portugal Romania Slovakia 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

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