# 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
Bupaq Multidose 0.3 mg/ml инжекционен разтвор за кучета и котки

**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:**
QN02AE01

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Bulgaria

**Available in:**
Bulgaria

**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:**
Bulgarian Food Safety Authority

**Authorisation number:**
0022-2142

**Date of authorisation status change:**
19/07/2016

**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

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.