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

Authorised

- Buprenorphine hydrochloride

Product identification

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

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:
Austria

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:
27/10/2011

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-01011

Date of authorisation status change:
27/10/2011

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

English (PDF)
Published on: 4/05/2023
Updated on: 3/05/2024
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Package Leaflet

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Labelling

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