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VETERGESIC MULTIDOSE, 0.3MG/ML, SOLUTION FOR INJECTION FOR DOGS AND CATS

Authorised

- Buprenorphine hydrochloride

Product identification

Medicine name:
VETERGESIC MULTIDOSE, 0.3MG/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:
Norway

Available in:
Norway

Package description:
10ml amber, Type I glass vial with chlorobutyl rubber stopper and a 20 mm aluminium collar with a flip-off cap

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
27/08/2009

Manufacturing sites for batch release:
Labiana Life Sciences S.A.

Responsible authority:

Norwegian Medical Products Agency

Authorisation number:
09-6502

Date of authorisation status change:
11/04/2013

Reference member state:
France

Procedure number:
FR/V/0368/001

Concerned member states:
Austria Belgium Czechia Denmark Finland Germany Hungary Iceland Ireland Luxembourg Netherlands Norway Poland Slovakia 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.

Package Leaflet

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

Combined File of all Documents

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