Morphasol 4 mg/ml solution for injection for dogs and cats

Authorised

- Butorphanol tartrate

Product identification

Medicine name:
Morphasol 4 mg/ml solution for injection for dogs and cats

Active substance:
Butorphanol tartrate

Target species:
Dog
Cat

Route of administration:
Intravenous use

Product details

Active substance and strength:
Butorphanol tartrate
5.83 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN02AF01

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

Authorisation status:
Valid

Authorised in:
Iceland

Available in:
Iceland

Package description:
Cardboard box with 1 glass vial (type I) of 10 ml with a grey butyl rubber stopper and an aluminium cap.
Cardboard box with 5 glass vial (type I) of 10 ml with grey butyl rubber stopper and an aluminium 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:
aniMedica GmbH

Marketing authorisation date:
1/12/2009

Manufacturing sites for batch release:
Industrial Veterinaria S.A.
aniMedica GmbH

Responsible authority:
Icelandic Medicines Agency

Authorisation number:
IS/2/09/015/01

Date of authorisation status change:
8/01/2015

Reference member state:
Ireland

Procedure number:
IE/V/0232/001

Concerned member states:
Austria Belgium Denmark Estonia Finland France Germany Greece Hungary Iceland Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Spain Sweden

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)
Published on: 28/09/2025
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Combined File of all Documents

Labelling

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Package Leaflet

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