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Myodine 25 mg/ml solution for injection for dogs and cats

Authorised

- Nandrolone laurate

Product identification

Medicine name:
Myodine 25 mg/ml solution for injection for dogs and cats

Active substance:
Nandrolone laurate

Target species:
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Nandrolone laurate
25.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA14AB01

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

Authorisation status:
Valid

Authorised in:
Spain

Package description:
Cardboard box with 1 clear type I glass vial containing 5 ml with a coated bromobutyl rubber stopper and aluminium cap.Pack size:Box with 1 vial of 5 ml
Cardboard box with 1 clear type II glass vial containing 10 ml with a coated bromobutyl rubber stopper and aluminium cap.Pack size:Box with 1 vial of 10 ml
Cardboard box with 1 clear type II glass vial containing 20 ml with a coated bromobutyl rubber stopper and aluminium cap.Pack size:Box with 1 vial of 20 ml
Cardboard box with 6 clear type I glass vials containing 5 ml with a coated bromobutyl rubber stopper and aluminium cap.Pack size:Multi-pack with 6 vials of 5 ml
Cardboard box with 6 clear type II glass vials containing 10 ml with a coated bromobutyl rubber stopper and aluminium cap.Pack size:Multi-pack with 6 vials of 10 ml
Cardboard box with 6 clear type II glass vials containing 20 ml with a coated bromobutyl rubber stopper and aluminium cap.Pack size:Multi-pack with 6 vials of 20 ml
Cardboard box with 10 clear type I glass vials containing 5 ml with a coated bromobutyl rubber stopper and aluminium cap.Pack size:Multi-pack with 10 vials of 5 ml
Cardboard box with 10 clear type II glass vials containing 10 ml with a coated bromobutyl rubber stopper and aluminium cap.Pack size:Multi-pack with 10 vials of 10 ml
Cardboard box with 10 clear type II glass vials containing 20 ml with a coated bromobutyl rubber stopper and aluminium cap.Pack size:Multi-pack with 10 vials of 20 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:
Le Vet. Beheer B.V.

Marketing authorisation date:
21/07/2017

Manufacturing sites for batch release:
Produlab Pharma B.V.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
3574 ESP

Date of authorisation status change:
7/03/2019

Reference member state:
Ireland

Procedure number:
IE/V/0358/001

Concerned member states:
Austria Belgium Croatia Cyprus Czechia Denmark Estonia Finland France Greece Hungary Iceland Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia 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: 20/09/2026
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Combined File of all Documents

Labelling

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Package Leaflet

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