Laurabolin 25 mg/ml solution for injection

Authorised

- Nandrolone laurate

Product identification

Medicine name:
Laurabolin 25 mg/ml solution for injection

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

Withdrawal period by route of administration:

Intramuscular use:

- **Dog**
- **Cat**

Subcutaneous use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA14AB01

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

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Authorisation status:
Valid

Authorised in:
Ireland

Package description:
Hydrolytic Type I glass vial with rubber stopper, closed with a colour coded aluminium cap. Each vial contains 10 ml solution.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet (Ireland) Limited

Marketing authorisation date:
1/10/1987

Manufacturing sites for batch release:
Intervet International GmbH
INTERVET INTERNATIONAL B.V.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10996/002/001

Date of authorisation status change:
1/10/1987

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

Documents

Summary of Product Characteristics
