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HEDYLON 25 mg TABLETS FOR DOGS

Authorised

- Prednisolone

Product identification

Medicine name:
HEDYLON 25 mg TABLETS FOR DOGS

Active substance:
Prednisolone

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Prednisolone
25.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB06

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

Authorisation status:
Valid

Authorised in:
Norway

Available in:
Norway

Package description:
cardboard box containing 10 tablets (1 blister)
cardboard box containing 30 tablets (3 blister)
cardboard box containing 50 tablets (5 blister)
cardboard box containing 100 tablets (10 blister)
cardboard box containing 250 tablets (25 blister)

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:
Industrial Veterinaria S.A.

Marketing authorisation date:
24/10/2019

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

aniMedica Herstellungs GmbH

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
19-12969

Date of authorisation status change:
24/10/2019

Reference member state:
Spain

Procedure number:
ES/V/0292/002

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Netherlands Norway Poland Portugal Romania Slovakia Slovenia 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: 20/10/2025
Updated on: 13/03/2026
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Combined File of all Documents

Package Leaflet

English (PDF)
Published on: 13/03/2026
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Labelling

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