HEDYLON 25 mg TABLETS FOR DOGS

Authorised

Prednisolone

Product identification

Medicine name:

HEDYLON 25 mg TABLETS FOR DOGS Hedylon, 25 mg tabletės šunims

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

Withdrawal period by route of administration:

Oral use:

. Dog

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB06

Legal status of supply:

This information is not available for this product.

Authorisation status:

Valid

Authorised in:

Lithuania

Package description:

cardboard box containing 250 tablets (25 blister)

cardboard box containing 100 tablets (10 blister)

cardboard box containing 50 tablets (5 blister)

cardboard box containing 30 tablets (3 blister)

cardboard box containing 10 tablets (1 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:

Livisto Int'l S.L.

Marketing authorisation date:

7/02/2019

Manufacturing sites for batch release:

Industrial Veterinaria S.A.

Animedica GmbH

Animedica Herstellungs GmbH
Responsible authority: State Food And Veterinary Service
Authorisation number: LT/2/19/2515/001-005
Date of authorisation status change: 7/02/2019
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
eu-PUAR-esv0292002-dcp-hedylon-25-mg-tablets-for-dogs-en.pdf

RV2515.pdf			
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