Source URL: https://medicines.health.europa.eu/veterinary/en/60000039793

Kelevo 400 µg tablets for dogs and cats

Authorised

• Levothyroxine sodium

Product identification

Medicine name:

Kelevo 400 μg δισκία για σκύλους και γάτες Kelevo 400 μg tablets for dogs and cats

Active substance:

Levothyroxine sodium

Target species:

Dog

Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Levothyroxine sodium 400.00 microgram(s) / 1.00 Tablet

Pharmaceutical form:

Withdrawal period by route of administration:

Oral use:

•

Dog

•

Cat

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH03AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Package description:

Aluminium - Aluminium blister each with 25 tablets. Cardboard box with 50 tablets

Aluminium - Aluminium blister each with 25 tablets. Cardboard box with 250 tablets

Aluminium - Aluminium blister each with 25 tablets. Cardboard box with 100 tablets

Aluminium - Aluminium blister each with 10 tablets. Cardboard box with 50 tablets.

Aluminium - Aluminium blister each with 10 tablets. Cardboard box with 250 tablets

Aluminium - Aluminium blister each with 10 tablets. Cardboard box with 100 tablets

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Industrial Veterinaria S.A.

Marketing authorisation date:

1/08/2021

Manufacturing sites for batch release:

aniMedica GmbH aniMedica Herstellungs GmbH Industrial Veterinaria S.A. Lelypharma B.V.

Responsible authority:

National Organization For Medicines

Authorisation number:

73683/02-08-2021/K-0245102

Date of authorisation status change:

1/08/2021

Reference member state:

Netherlands

Procedure number:

NL/V/0349/002

Concerned member states:

Austria Estonia Germany Greece Hungary Ireland Italy Latvia Lithuania Poland Portugal Slovenia Spain

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

Documents

Combined File of all Documents

Summary of Product Characteristics

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

Package Leaflet

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