Kelevo 200 µg tablets for dogs and cats

Authorised

- Anhydrous levothyroxine sodium

Product identification

Medicine name:
Kelevo 200 µg tablets for dogs and cats
KELEVO 200 µg COMPRIMIDOS PARA PERROS Y GATOS

Active substance:
Anhydrous levothyroxine sodium

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Anhydrous levothyroxine sodium
200.00 microgram(s) / 1.00 Piece

Pharmaceutical form:
Tablet

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:

Spain

Available in:

Spain

Package description:

PVC / PE / PVDC - Aluminium blister each with 25 tablets. Cardboard box with 50 tablets.
PVC / PE / PVDC - Aluminium blister each with 25 tablets. Cardboard box with 250 tablets.
PVC / PE / PVDC - Aluminium blister each with 25 tablets. Cardboard box with 100 tablets.
PVC / PE / PVDC - Aluminium blister each with 10 tablets. Cardboard box with 50 tablets.
PVC / PE / PVDC - Aluminium blister each with 10 tablets. Cardboard box with 250 tablets.
PVC / PE / PVDC - Aluminium blister each with 10 tablets. Cardboard box with 100 tablets.

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:
7/04/2021

Manufacturing sites for batch release:
Animedica GmbH
Animedica Herstellungs GmbH
Industrial Veterinaria S.A.
Lelypharma B.V.

Responsible authority:
Spanish Agency For Medicines And Health Products

Authorisation number:
3993 ESP

Date of authorisation status change:
8/04/2021

Reference member state:
Netherlands

Procedure number:
NL/V/0349/001

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

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.

Labelling

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

Combined File of all Documents
