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Kelevo 800 µg tablets for dogs

Authorised

- Levothyroxine sodium

Product identification

Medicine name:
Kelevo 800 µg tablets for dogs

Active substance:
Levothyroxine sodium

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Levothyroxine sodium
800.00 microgram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH03AA01

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

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
Aluminium - Aluminium blister each with 25 tablets. Cardboard box with 100 tablets
Aluminium - Aluminium blister each with 25 tablets. Cardboard box with 250 tablets
Aluminium - Aluminium blister each with 10 tablets. Cardboard box with 250 tablets
Aluminium - Aluminium blister each with 10 tablets. Cardboard box with 50 tablets.
Aluminium - Aluminium blister each with 10 tablets. Cardboard box with 100 tablets
Aluminium - Aluminium blister each with 25 tablets. Cardboard box with 50 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:
15/04/2021

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

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402765.00.00

Date of authorisation status change:
15/04/2021

Reference member state:
Netherlands

Procedure number:
NL/V/0349/003

Concerned member states:
Austria Estonia Germany Greece Hungary Ireland Italy Latvia Lithuania Poland Portugal Slovenia Spain

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Documents

Combined File of all Documents

English (PDF)
Published on: 11/11/2025
Updated on: 14/03/2026
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