

Forthyron 800 Microgram Flavoured Tablet

Authorised

- Levothyroxine sodium

Product identification

Medicine name:

Forthyron 800 Microgram Flavoured Tablet

Forthyron flavoured 800 µg δισκία για σκύλους

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:

Cyprus

Package description:

10 tablets in a blister [Aluminium (20micrometre) - PVC_PE_PVDC (250_30_90) white],
25 blisters per carton

10 tablets in a blister [Aluminium (20micrometre) - PVC_PE_PVDC (250_30_90) white],
5 blisters per carton

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:

Eurovet Animal Health B.V.

Marketing authorisation date:

27/09/2018

Manufacturing sites for batch release:

Dales Pharmaceuticals Limited

Eurovet Animal Health B.V.

Genera d.d.

Responsible authority:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

CY00707V

Date of authorisation status change:

27/09/2018

Reference member state:

Netherlands

Procedure number:

NL/V/0286/004

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland
France Germany Greece Hungary Ireland Italy Latvia Lithuania
Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain
Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)

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