Forthyron 400 Microgram Flavoured Tablet

Authorised

- Levothyroxine sodium

Product identification

Medicine name:
Forthyron 400 Microgram Flavoured Tablet

Active substance:
Levothyroxine sodium

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Levothyroxine sodium
400.00 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:
Netherlands

Available in:
Netherlands

Package description:
10 tablets in a blister [Aluminium (20micrometre) - PVC_PE_PVDC (250_30_90) white], 5 blisters per carton
10 tablets in a blister [Aluminium (20micrometre) - PVC_PE_PVDC (250_30_90) white], 25 blisters per carton

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:
Eurovet Animal Health B.V.

Marketing authorisation date:
29/11/2011

Manufacturing sites for batch release:
Dales Pharmaceuticals Limited
Eurovet Animal Health B.V.
Genera d.d.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 108734

Date of authorisation status change:
24/01/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0286/002

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)
Published on: 20/06/2022
Updated on: 21/06/2022
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