Forthyron 600 Microgram Flavoured Tablet

Authorised

- Levothyroxine sodium

Product identification

Medicine name:
Forthyron 600 Microgram Flavoured Tablet
Forthyron flavour, 600 µg tabletid koertele

Active substance:
Levothyroxine sodium

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Levothyroxine sodium
600.00 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:
Estonia

Available in:
Estonia

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

Marketing authorisation holder:
Eurovet Animal Health B.V.

Marketing authorisation date:
4/09/2018

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

Responsible authority:

State Agency Of Medicines

Authorisation number:

2107

Date of authorisation status change:

4/09/2018

Reference member state:

Netherlands

Procedure number:

NL/V/0286/003

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland 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

Summary of Product Characteristics

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

Combined File of all Documents

108735 - par.pdf