Soloxine 0.1 mg Tablet

Authorised

- Levothyroxine sodium

Product identification

Medicine name:
Soloxine 0.1 mg Tablet

Active substance:
Levothyroxine sodium

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Levothyroxine sodium
0.10 milligram(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:
Ireland

Package description:
High-density, brown, polyethylene bottles containing 250 tablets, hermetically sealed and closed with childproof screwcap

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Virbac

Marketing authorisation date:
19/01/2007

Manufacturing sites for batch release:
Virbac

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10988/069/001

Date of authorisation status change:
19/01/2007

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

Documents

Summary of Product Characteristics