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Soloxine 0.5 mg Tablet

Authorised

- Levothyroxine sodium

Product identification

Medicine name:

Soloxine 0.5 mg Tablet

Active substance:

Levothyroxine sodium

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Levothyroxine sodium
0.50 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:

VPA10988/069/004

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