

Synulox 50 mg tablets for dogs and cats

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:

Synulox 50 mg tablets for dogs and cats

Active substance:

Potassium clavulanate

Amoxicillin trihydrate

Target species:

Dog

Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Potassium clavulanate

11.91 milligram(s) / 1.00 Tablet

Amoxicillin trihydrate
45.91 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CR02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Finland

Available in:

Finland

Package description:

(ID3) 500 Tablet: Box (cardboard) with 50 Blister (aluminium; low-density polyethylene) each with 10 Tablet

(ID2) 100 Tablet: Box (cardboard) with 10 Blister (aluminium; low-density polyethylene) each with 10 Tablet

(ID1) 10 Tablet: Box (cardboard) with 1 Blister (aluminium; low-density polyethylene) with 10 Tablet

(ID4) 250 Tablet: Box (cardboard) with 25 Blister (aluminium; low-density polyethylene) each with 10 Tablet

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Zoetis Animal Health ApS

Marketing authorisation date:

23/10/1990

Manufacturing sites for batch release:

Haupt Pharma Latina S.r.l.

Responsible authority:

Finnish Medicines Agency

Authorisation number:

10314

Date of authorisation status change:

23/10/1990

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

Documents

Package Leaflet

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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