

Synulox 500 mg tablets for dogs

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:

Synulox 500 mg Comprimés Appétents

Synulox 500 mg tablets for dogs

Active substance:

Potassium clavulanate

Amoxicillin trihydrate

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Potassium clavulanate

119.14 milligram(s) / 1.00 Tablet

Amoxicillin trihydrate

459.10 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:

Luxembourg

Package description:

(ID3) 20 Tablet: Box (board) with 10 Blister (aluminium; low-density polyethylene) each with 2 Tablet

(ID2) 100 Tablet: Box (board) with 50 Blister (aluminium; low-density polyethylene) each with 2 Tablet

(ID1) 10 Tablet: Box (board) with 5 Blister (aluminium; low-density polyethylene) each with 2 Tablet

(ID4) 200 Tablet: Box (board) with 100 Blister (aluminium; low-density polyethylene) each with 2 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 Belgium

Marketing authorisation date:

25/11/1998

Manufacturing sites for batch release:

Haupt Pharma Latina S.r.l.

Responsible authority:

Ministry Of Health And Social Security

Authorisation number:

V 087/98/11/0627

Date of authorisation status change:

23/10/2007

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