

Synulox 40 mg/ml + 10 mg/ml powder for oral suspension for dogs and cats

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:

Synulox 40 mg/ml + 10 mg/ml powder for oral suspension 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

193.00 milligram(s) / 1.70 gram(s)

Amoxicillin trihydrate

743.80 milligram(s) / 1.70 gram(s)

Pharmaceutical form:

Powder for oral suspension

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) 1.7 gram(s): Box (board) with 1 Bottle (clear glass) with 1.7 gram(s), closed with (Metall)

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:

20/12/2004

Manufacturing sites for batch release:

Haupt Pharma Latina S.r.l.

Responsible authority:

Ministry Of Health And Social Security

Authorisation number:

V 087/04/12/0600

Date of authorisation status change:

9/11/2009

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