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 POLVO PARA GOTAS ORALES EN SUSPENSION
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:
Spain

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 Spain S.L.

Marketing authorisation date:
11/03/1987

Manufacturing sites for batch release:

Haupt Pharma Latina S.r.l.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
2574 ESP

Date of authorisation status change:
27/06/2012

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

Documents

Summary of Product Characteristics

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Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents