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

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Combined File of all Documents