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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:
Germany

Available in:
Germany

Package description:
Available only in German

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 Deutschland GmbH

Marketing authorisation date:
7/04/2004

Manufacturing sites for batch release:

Haupt Pharma Latina S.r.l.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
400602.00.01

Date of authorisation status change:
9/07/2009

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

Documents

Combined File of all Documents

English (PDF)
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Updated on: 14/03/2026
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