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Synulox 500 mg tablets for dogs

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:
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:
Germany

Available in:
Germany

Package description:
(ID4) 200 Tablet: Box (cardboard) with 100 Blister (aluminium; low-density polyethylene) each with 2 Tablet
(ID4) 200 Tablet: Box (cardboard) with 100 Blister (aluminium; low-density polyethylene) each with 2 Tablet
(ID4) 200 Tablet: Box (cardboard) with 100 Blister (aluminium; low-density polyethylene) each with 2 Tablet
(ID4) 200 Tablet: Box (cardboard) 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 Deutschland GmbH

Marketing authorisation date:

5/12/2003

Manufacturing sites for batch release:
Haupt Pharma Latina S.r.l.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
400602.02.00

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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