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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:
Norway

Available in:
Norway

Package description:
(ID3) 20 Tablet: Box (cardboard) with 10 Blister (aluminium; low-density polyethylene) each with 2 Tablet
(ID2) 100 Tablet: Box (cardboard) with 50 Blister (aluminium; low-density polyethylene) each with 2 Tablet
(ID1) 10 Tablet: Box (cardboard) with 5 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 Animal Health ApS

Marketing authorisation date:

7/03/1997

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

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
1995-01061

Date of authorisation status change:
3/07/2007

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

Documents

Package Leaflet

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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