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:
Greece

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 Hellas S.A.

Marketing authorisation date:
9/05/2004

Manufacturing sites for batch release:

Haupt Pharma Latina S.r.l.

Responsible authority:
National Organization For Medicines

Authorisation number:
97735/19-10-2021/K-0077406

Date of authorisation status change:
18/10/2021

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

Documents

Combined File of all Documents