Synuclav 200 mg / 50 mg Tablets for Dogs

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:
Synuclav 200 mg / 50 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
59.50 milligram(s) / 1.00 Tablet

Amoxicillin trihydrate

229.50 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:
France

Available in:
France

Package description:
box containing 20 tablets (4 blister)
box containing 50 tablets (10 blister)
box containing 250 tablets (50 blister)
bottle containing 100 tablets
bottle containing 250 tablets
box containing 100 tablets (20 blister)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:

4/06/2004

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Laboratories (Ireland) Limited

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/1889296 5/2004

Date of authorisation status change:
7/12/2019

Reference member state:
Spain

Procedure number:
ES/V/0349/002

Concerned member states:
Belgium France Iceland Italy Portugal United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 22/10/2025
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Package Leaflet