Synuclav Suspension for Injection

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:
Synuclav Suspension for Injection
CLAVOBAY SUSPENSION INJECTABLE

Active substance:
Potassium clavulanate
Amoxicillin trihydrate

Target species:
Cattle
Dog

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Potassium clavulanate
35.00 milligram(s) / 1.00 millilitre(s)

Amoxicillin trihydrate
140.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 42 day

- **Cattle**
  - Milk. 60 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CR02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription except for some pack sizes

Authorisation status:
Valid

Authorised in:
France

Available in:
France

Package description:
box containing 1 vial of 100 ml
box containing 1 vial of 50 ml

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:
11/06/2004

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/7057150 1/2004

Date of authorisation status change:
11/06/2009

Reference member state:
Spain

Procedure number:
ES/V/0354/001

Concerned member states:
Belgium France Iceland Italy Portugal

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 23/03/2023
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Combined File of all Documents