

Synoclav Suspension for Injection

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:

Synoclav Suspension for Injection
NOROCLAV 175 mg SUSPENSION INYECTABLE

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

Authorisation status:

Valid

Authorised in:

Spain

Available in:

Spain

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:

28/07/2004

Manufacturing sites for batch release:

Norbrook Laboratories Limited

Norbrook Laboratories (Ireland) Limited

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

1582 ESP

Date of authorisation status change:

26/02/2018

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

Combined File of all Documents

Summary of Product Characteristics

English (PDF)

Published on: 23/03/2023

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Package Leaflet

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Labelling

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