

CLAMOX RTU 140/35 mg/ml suspensie injectabilă pentru bovine, porci, câini și pisici

Authorised

- Potassium clavulanate
- Amoxicillin

Product identification

Medicine name:

CLAMOX RTU 140/35 mg/ml suspensie injectabilă pentru bovine, porci, câini și pisici

Active substance:

Potassium clavulanate

Amoxicillin

Target species:

Cattle

Pig

Dog

Cat

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Potassium clavulanate

35.00 milligram(s) / 1.00 millilitre(s)

Amoxicillin

140.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Powder for solution for injection

Withdrawal period by route of administration:

Intramuscular use:

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Cattle

- Meat and offal. 42 day

- Milk. 60 hour

În cazul terapiei combinate, laptele va fi dat în consum după 60 ore de la ultimul tratament (după 5 mulsori, în cazul în care vacile sunt mulse de două ori pe zi).

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Pig

- Meat and offal. 31 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CR02

Legal status of supply:

This information is not available for this product.

Authorisation status:

Valid

Authorised in:

Romania

Package description:

Available only in [Romanian](#)

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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:

Crida Pharm S.R.L.

Marketing authorisation date:

27/06/2018

Manufacturing sites for batch release:

Crida Pharm S.R.L.

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

230110

Date of authorisation status change:

18/03/2026

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

Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.