Dokamox 100 mg/g

Not authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Dokamox 100 mg/g

Active substance:
Amoxicillin trihydrate

Target species:
Pig

Route of administration:
Oral use

Product details

Active substance and strength:
Amoxicillin trihydrate
114.79 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Premix for medicated feeding stuff

Withdrawal period by route of administration:

Oral use:

- **Pig**
  - Meat and offal. 5 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Surrendered

Authorised in:
Italy

Package description:
25 kg low density polyethylene / paper / paper bag
10 kg low density polyethylene / paper / paper bag

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Ceva Salute Animale S.p.A.

Marketing authorisation date:
2/09/2010

Manufacturing sites for batch release:
CEVA Santé Animale

Responsible authority:
Ministry Of Health

Authorisation number:
104080

Date of authorisation status change:
27/01/2022

Reference member state:
Belgium

Procedure number:
BE/V/0047/001

Generic of:
600000040301

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

Documents

Combined File of all Documents

English (PDF)
Published on: 13/03/2026
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Package Leaflet

Summary of Product Characteristics