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Amatib 800 mg/g oral powder for pigs and chickens

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Amatib 800 mg/g oral powder for pigs and chickens

Active substance:
Amoxicillin trihydrate

Target species:
Chicken
Pig

Route of administration:
Oral use

Product details

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

Pharmaceutical form:
Powder for use in drinking water

Withdrawal period by route of administration:

Oral use:

- **Chicken**
  - Eggs. no withdrawal period

Not authorised for use in laying birds producing eggs for human consumption. Do not use within 3 weeks of onset of lay.

  - Meat and offal. 1 day

- **Pig**
  - Meat and offal. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

Thermosealed bags of PET/Al/PE containing 1000 g powder.

Thermosealed bags of PET/Al/PE containing 500 g powder.

Thermosealed bags of PET/Al/PE containing 250 g powder.

Thermosealed bags of PET/Al/PE containing 100 g powder.

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:
KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:
3/08/2015

Manufacturing sites for batch release:
TAD Pharma GmbH
KRKA tovarna zdravil d.d. Novo mesto

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

Authorisation number:
FR/V/2281788 2/2015

Date of authorisation status change:
31/07/2020

Reference member state:
Ireland

Procedure number:
IE/V/0346/001

Concerned member states:
Belgium Bulgaria Croatia Czechia Estonia France Germany Hungary Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Package Leaflet and Labelling

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

Summary of Product Characteristics

English (PDF)
Published on: 3/05/2024
Updated on: 14/03/2026
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