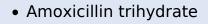
Huvamox 800 mg/g powder for use in drinking water for chickens, turkeys, ducks and pigs

Authorised



Product identification

Medicine name:

Huvamox 800 mg/g powder for use in drinking water for chickens, turkeys, ducks and pigs Huvamox 800 mg/g Proszek do podania w wodzie do picia

Active substance:

Amoxicillin trihydrate

Target species:

Turkey Chicken Duck Pig

Route of administration:

In drinking water 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:

In drinking water use:

- Turkey
 - Meat and offal. 5 day
- . Chicken
 - Meat and offal. 1 day
- Duck
 - Meat and offal. 9 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

Authorisation status:

Valid

Authorised in:

Poland

Package description:

Zipped 1 kg bag made of low density polyethylene/aluminium/polyethylene terephthalate.

Zipped 500 g bag made of low density polyethylene/aluminium/polyethylene terephthalate.

Thermo-sealed 100 g bag made of low density polyethylene/aluminium/polyethylene terephthalate.

100 g jar made of high density polyethylene closed with a seal made of low density polyethylene/ polyethylene terephthalate/aluminium and a screw cap made of polypropylene.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

HuVepharma

Marketing authorisation date:

26/05/2021

Manufacturing sites for batch release:

Huvepharma

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

3096

Reference member state:

Ireland

Procedure number:

IE/V/0642/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia France Germany Greece Hungary Italy Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF) Published on: 11/02/2022 Download

Labelling

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Package Leaflet

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

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