

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

Authorised

- Amoxicillin trihydrate

## Product identification

### Medicine name:

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

HUVAMOX 800 mg/g κόνις για χορήγηση με πόσιμο νερό για ορνίθια, ινδόρνιθες, πάπιες και χοίρους

### 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)

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**Pharmaceutical form:**

Powder for use in drinking water

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**Withdrawal period by route of administration:****In drinking water use:**

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**Turkey**

- Meat and offal. 5 day

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**Chicken**

- Meat and offal. 1 day

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**Duck**

- Meat and offal. 9 day

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**Pig**

- Meat and offal. 2 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01CA04

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Cyprus

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**Available in:**

Cyprus

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**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/ polyethyleneterephthalate/aluminium and a screw cap made of polypropylene.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

HuVepharma

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**Marketing authorisation date:**

11/04/2021

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**Manufacturing sites for batch release:**

Huvepharma S.A.

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**Responsible authority:**

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

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**Authorisation number:**

CY00827V

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**Date of authorisation status change:**

11/04/2021

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0642/001

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**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)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

Combined File of all Documents

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