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# 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

**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**
  - Eggs. no withdrawal period

Not authorised for use in laying birds producing eggs for human consumption. Do not use within 4 weeks of the start of the laying period.

  - Meat and offal. 5 day

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

Not authorised for use in laying birds producing eggs for human consumption. Do not use within 4 weeks of the start of the laying period.

  - Meat and offal. 1 day

- **Duck**
  - Eggs. no withdrawal period

Not authorised for use in laying birds producing eggs for human consumption. Do not use within 4 weeks of the start of the laying period.

  - 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:**
Romania

**Available in:**
Romania

**Package description:**
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.
Thermo-sealed 100 g bag made of low density polyethylene/aluminium/polyethylene terephthalate.
Zipped 500 g bag made of low density polyethylene/aluminium/polyethylene terephthalate.
Zipped 1 kg bag made of low density polyethylene/aluminium/polyethylene terephthalate.

## 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:**
2/09/2021

**Manufacturing sites for batch release:**
Huvepharma S.A.

**Responsible authority:**
Institute For Control Of Biological Products And Veterinary Medicines

**Authorisation number:**
260123

**Date of authorisation status change:**
1/09/2026

**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)

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