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 697 MG/G POUDRE POUR ADMINISTRATION DANS L'EAU DE BOISSON POUR POULES, DINDES, CANARDS ET PORCS

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 except for some pack sizes

Authorisation status:
Valid

Authorised in:
France

Available in:
France

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.

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:
3/03/2021

Manufacturing sites for batch release:
Huvepharma S.A.

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

Authorisation number:
FR/V/5558375 2/2021

Date of authorisation status change:
3/03/2021

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 www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)
Published on: 3/05/2024
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Package Leaflet and Labelling

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