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

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Awazom 800 mg/g Powder for Use in Drinking Water for Chickens, Ducks and Turkeys
Awazom 800 mg/g powder for use in drinking water for chickens, ducks and turkeys

Active substance:
Amoxicillin trihydrate

Target species:
Chicken
Duck
Turkey

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:

- **Chicken**
  - Meat and offal. 1 day

Ni za uporabo pri pticah, ki proizvajajo jajca za prehrano ljudi. Ne uporabite v obdobju 3 tednov pred začetkom obdobja nesnosti.

- **Duck**
  - Meat and offal. 9 day

Ni za uporabo pri pticah, ki proizvajajo jajca za prehrano ljudi. Ne uporabite v obdobju 3 tednov pred začetkom obdobja nesnosti.

- **Turkey**
  - Meat and offal. 5 day

Ni za uporabo pri pticah, ki proizvajajo jajca za prehrano ljudi. Ne uporabite v obdobju 3 tednov pred začetkom obdobja nesnosti.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
United Kingdom (Northern Ireland)

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

Marketing authorisation holder:
KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:
9/04/2019

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

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 01656/3071

Date of authorisation status change:
9/04/2024

Reference member state:
Slovenia

Procedure number:
SI/V/0002/001

Concerned member states:

United Kingdom (Northern Ireland)

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