TUDOMAX 10 MG/G, POWDER FOR USE IN DRINKING WATER/MILK

Authorised

- Bromhexine hydrochloride

Product identification

Medicine name:
TUDOMAX 10 MG/G, POWDER FOR USE IN DRINKING WATER/MILK

Active substance:
Bromhexine hydrochloride

Target species:
Turkey
Pig
Cattle (calf)
Duck
Chicken
Chicken (broiler)

Route of administration:
Oral use

Product details

Active substance and strength:
Bromhexine hydrochloride

10.98 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for use in drinking water/milk

Withdrawal period by route of administration:
Oral use:

- **Turkey**
  - Meat and offal. 0 day
  - Eggs. no withdrawal period

Not for use in birds producing eggs for consumption, during and 4 weeks before the laying phase.

- **Pig**
  - Meat and offal. 0 day

- **Cattle (calf)**
  - Meat and offal. 2 day
  - Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

- **Duck**
  - Meat and offal. 0 day
  - Eggs. no withdrawal period

Not for use in birds producing eggs for consumption, during and 4 weeks before the laying phase.

- **Chicken**

- Eggs. no withdrawal period

Not for use in birds producing eggs for consumption, during and 4 weeks before the laying phase.

- **Chicken (broiler)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QR05CB02

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

Authorisation status:
Valid

Authorised in:
Romania

Available in:
Romania

Package description:
Bag of 500 g
Bag of 1 kg

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:
S P Veterinaria S.A.

Marketing authorisation date:
20/02/2017

Manufacturing sites for batch release:
S P Veterinaria S.A.

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

Authorisation number:
220030

Date of authorisation status change:
28/05/2025

Reference member state:
France

Procedure number:
FR/V/0295/001

Concerned member states:
Bulgaria Cyprus Greece Hungary Ireland Italy Malta Poland Portugal Romania Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)
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