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
TUDOMAX 10 mg/g POLVO PARA ADMINISTRACIÓN EN AGUA DE BEBIDA O EN LECHE

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:
Spain

Available in:
Spain

Package description:
Available only in French
Available only in French

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:
6/02/2017

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

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
3517 ESP

Date of authorisation status change:
8/02/2017

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

Summary of Product Characteristics

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Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.