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DOPHEXINE 20 MG/G POWDER FOR USE IN DRINKING WATER/MILK

Authorised

- Bromhexine hydrochloride

Product identification

Medicine name:
DOPHEXINE 20 MG/G POWDER FOR USE IN DRINKING WATER/MILK

Active substance:
Bromhexine hydrochloride

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

Route of administration:
Oral use

Product details

Active substance and strength:
Bromhexine hydrochloride
20.00 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 human consumption, during and 4 weeks before the laying period.

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

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

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

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

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

- **Chicken (broiler)**
  - Meat and offal. 0 day
- **Cattle**
  - Meat and offal. 2 day
  - Milk. no withdrawal period

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QR05CB02

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

Authorisation status:
Valid

Authorised in:
Lithuania

Available in:
Lithuania

Package description:
Composite can of 1 kg
Bucket of 5 kg
Bucket of 2,5 kg
Bucket of 1 kg
Securitainer 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:
Dopharma Research B.V.

Marketing authorisation date:
16/08/2020

Manufacturing sites for batch release:
Dopharma B.V.

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/20/2613/001-005

Date of authorisation status change:
23/11/2025

Reference member state:
France

Procedure number:
FR/V/0391/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Finland Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Spain Sweden

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Documents

Combined File of all Documents

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