EXFLOW 10 MG/G POWDER FOR USE IN DRINKING WATER FOR CATTLE (CALVES), PIGS, CHICKENS, TURKEYS AND DUCKS

Not authorised

- Bromhexine hydrochloride

Product identification

Medicine name:
EXFLOW 10 MG/G POWDER FOR USE IN DRINKING WATER FOR CATTLE (CALVES), PIGS, CHICKENS, TURKEYS AND DUCKS
Exflow 10 mg / g prášok na použitie v pitnej vode pre hovädzí dobytok (teľatá), ošípané, kurčatá, morky a kačice

Active substance:
Bromhexine hydrochloride

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

Route of administration:
Oral use

Product details

Active substance and strength:
Bromhexine hydrochloride
10.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for use in drinking water

Withdrawal period by route of administration:
Oral use:

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

Do not 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 permitted for use in cows producing milk for human consumption.

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

- Eggs. no withdrawal period

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

- **Chicken (broiler)**
  - Meat and offal. 0 day
  - Eggs. no withdrawal period

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QR05CB02

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

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Authorisation status:
Surrendered

Authorised in:
Slovakia

Package description:
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Animal Health Slovakia s.r.o.

Marketing authorisation date:
17/12/2015

Manufacturing sites for batch release:
Ceva Sante Animale
Laboratoires Biove

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/079/DC/15-S

Date of authorisation status change:
27/03/2024

Reference member state:
France

Procedure number:
FR/V/0285/001

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

Documents

eu-puar-frv0285001-mr-rpe184-en.pdf