

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:

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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.

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Pig

- Meat and offal. 0 day

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Cattle (calf)

- Meat and offal. 2 day
- Milk. no withdrawal period

Not permitted for use in cows producing milk for human consumption.

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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.

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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

Authorisation status:

Surrendered

Authorised in:

Slovakia

Package description:

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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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 Santé 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

Source URL: <https://medicines.health.europa.eu/veterinary/600000030183>