

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

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:

Revoked

Authorised in:

Portugal

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 Saude Animal Produtos Farmaceuticos E Imunologicos Lda.

Marketing authorisation date:

10/08/2015

Manufacturing sites for batch release:

Ceva Sante Animale
Laboratoires Biove

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

941/01/15RDVPT

Date of authorisation status change:

1/08/2024

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