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BROMHEX-AIR BASIC ORAL POWDER FOR CATTLE, PIGS, CHICKENS, TURKEYS AND DUCKS

Authorised

- Bromhexine hydrochloride

Product identification

Medicine name:
BROMHEX-AIR BASIC ORAL POWDER FOR CATTLE, PIGS, CHICKENS, TURKEYS AND DUCKS

Active substance:
Bromhexine hydrochloride

Target species:
Turkey
Pig
Duck
Chicken (broiler)
Cattle

Route of administration:
Oral use

Product details

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

Pharmaceutical form:
Oral powder

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. Do not use within 4 weeks before the start of 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. Do not use within 4 weeks before the start of the laying period.

- **Chicken (broiler)**
  - Meat and offal. 0 day

- Eggs. no withdrawal period

Not for use in birds producing eggs for human consumption. Do not use within 4 weeks before the start of the laying period.

- **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:
Luxembourg

Package description:
1 kg Bag
Box of 10 containers of 100 g
100 g container

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:

Pharmanovo Veterinaerarzneimittel GmbH

Marketing authorisation date:
6/09/2021

Manufacturing sites for batch release:
Animed Service AG

Responsible authority:
Ministry Of Health And Social Security

Authorisation number:
V/010/21/09/1850

Date of authorisation status change:
6/09/2021

Reference member state:
France

Procedure number:
FR/V/0426/001

Concerned member states:
Austria Belgium Luxembourg Netherlands

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

Documents

Summary of Product Characteristics

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Combined File of all Documents

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