

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

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

eu-puar-frv0426001-mr-rpe641-en.pdf