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

Authorised

- Bromhexine hydrochloride

Product identification

Medicine name:
BROMHEX-AIR FORTE 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
25.00 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 during and 4 weeks before the laying period.

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

- **Duck**
  - Eggs. no withdrawal period

Not for use in birds producing eggs for human consumption during and 4 weeks before the laying period.

  - Meat and offal. 0 day

- **Chicken (broiler)**
  - Eggs. no withdrawal period

Not for use in birds producing eggs for human consumption during and 4 weeks before the laying period.

- Meat and offal. 0 day

- **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:
France

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

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Pharmanovo Veterinaerarzneimittel GmbH

Marketing authorisation date:

18/05/2021

Manufacturing sites for batch release:
Animed Service AG

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/1192100 0/2021

Date of authorisation status change:
18/05/2021

Reference member state:
France

Procedure number:
FR/V/0426/002

Concerned member states:
Germany Hungary Ireland Italy Poland United Kingdom (Northern Ireland)

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

Package Leaflet and Labelling

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

eu-puar-frv0426002-mr-rpe642-en.pdf