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
Bromhex-Air forte 25 mg/g proszek doustny dla bydła świń, kur, indyków i kaczek

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:
Poland

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:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Pharmanovo Veterinaerarzneimittel GmbH

Marketing authorisation date:

29/04/2022

Manufacturing sites for batch release:
Animed Service AG

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3177

Date of authorisation status change:
29/04/2022

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

Package Leaflet

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Labelling

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Summary of Product Characteristics

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