

Bromhex-Air Forte 25 mg/g Oral Powder for Cattle, Pigs, Chickens, Turkeys and Ducks

Authorised

- Bromhexine hydrochloride

Product identification

Medicine name:

Bromhex-Air Forte 25 mg/g 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:

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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.

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Pig

- Meat and offal. 0 day

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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

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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

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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:

United Kingdom (Northern Ireland)

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 Veterinaerarztneimittel GmbH

Marketing authorisation date:

23/07/2021

Manufacturing sites for batch release:

Animed Service AG

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 52644/4000

Date of authorisation status change:

12/03/2024

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

Combined File of all Documents

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