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BRONCHORYL SOLUTION BUVABLE

Authorised

- Kalium stibyltartaricum C5
- Pulsatilla vulgaris C5
- Lobaria pulmonaria C4
- RUMEX CRISPUS C5
- DROSERA C3
- Atropa bella-donna C5
- ACONITUM NAPELLUS C5
- IPECA C3
- BRYONIA C5

Product identification

Medicine name:
BRONCHORYL SOLUTION BUVABLE

Active substance:
Kalium stibyltartaricum C5

Pulsatilla vulgaris C5

Lobaria pulmonaria C4

RUMEX CRISPUS C5

DROSERA C3

Atropa bella-donna C5

ACONITUM NAPELLUS C5

IPECA C3

BRYONIA C5

Target species:
Cattle
Pig
Rabbit
Horse
Sheep
Goat

Route of administration:
Oral use

Product details

Active substance and strength:
Kalium stibyltartaricum C5
108.80 milligram(s) / 1.00 millilitre(s)

Pulsatilla vulgaris C5
108.80 milligram(s) / 1.00 millilitre(s)

Lobaria pulmonaria C4
108.80 milligram(s) / 1.00 millilitre(s)

RUMEX CRISPUS C5
108.80 milligram(s) / 1.00 millilitre(s)

DROSERA C3
108.80 milligram(s) / 1.00 millilitre(s)

Atropa bella-donna C5
108.80 milligram(s) / 1.00 millilitre(s)

ACONITUM NAPELLUS C5
108.80 milligram(s) / 1.00 millilitre(s)

IPECA C3
108.80 milligram(s) / 1.00 millilitre(s)

BRYONIA C5
108.80 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral solution

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig**
  - All relevant tissues. 0 day
- **Rabbit**
  - All relevant tissues. 0 day
- **Horse**
  - All relevant tissues. 0 day
- **Sheep**
  - All relevant tissues. 0 day
  - Milk. 0 day
- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QV03AX

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:
France

Available in:
France

Package description:
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Homeopathic medicinal products (Article 19 of Directive No 2001/82/EC)

Marketing authorisation holder:
Boiron

Marketing authorisation date:
15/06/2012

Manufacturing sites for batch release:
Boiron

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

Authorisation number:
FR/V/4705529 0/2012

Date of authorisation status change:
31/05/2017

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

Documents

Summary of Product Characteristics

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Package Leaflet and Labelling

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