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ABCEDYL GA SOLUTION BUVABLE

Authorised

- PYROGENIUM C7
- Virola sebifera C3
- Atropa bella-donna C5
- ECHINACEA C3
- Calcium sulfuricum C7
- ACIDUM SILICICUM C7
- HEPAR SULFURIS C7

Product identification

Medicine name:
ABCEDYL GA SOLUTION BUVABLE

Active substance:
PYROGENIUM C7

Virola sebifera C3

Atropa bella-donna C5

ECHINACEA C3

Calcium sulfuricum C7

ACIDUM SILICICUM C7

HEPAR SULFURIS C7

Target species:
Cattle
Pig
Rabbit

Horse
Sheep
Goat
Poultry

Route of administration:
Oral use

Product details

Active substance and strength:
PYROGENIUM C7
139.90 milligram(s) / 1.00 millilitre(s)

Virola sebifera C3
139.90 milligram(s) / 1.00 millilitre(s)

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

ECHINACEA C3
139.90 milligram(s) / 1.00 millilitre(s)

Calcium sulfuricum C7
139.90 milligram(s) / 1.00 millilitre(s)

ACIDUM SILICICUM C7
139.90 milligram(s) / 1.00 millilitre(s)

HEPAR SULFURIS C7
139.90 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Withdrawal period by route of administration:
Oral use:

- **Cattle**
  - All relevant tissues. 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
- **Goat**
  - All relevant tissues. 0 day
- **Poultry**
  - All relevant tissues. 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:
28/06/2012

Manufacturing sites for batch release:
Boiron

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

Authorisation number:
FR/V/2653261 5/2012

Date of authorisation status change:
21/06/2017

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.