

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

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

Package Leaflet and Labelling

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