DRAINOSYL SOLUTION BUVABLE

Authorised

- TARAXACUM OFFICINALE D3
- BERBERIS VULGARIS C3
- SOLIDAGO VIRGAUREA C3
- CHINA D3
- Hydrastis canadensis C3
- Cynara scolymus D3
- SILYBUM MARIANUM C3
- Chelidonium majus C5

Product identification

Medicine name:
DRAINOSYL SOLUTION BUVABLE

Active substance:
TARAXACUM OFFICINALE D3

BERBERIS VULGARIS C3

SOLIDAGO VIRGAUREA C3

CHINA D3

Hydrastis canadensis C3

Cynara scolymus D3

SILYBUM MARIANUM C3

Chelidonium majus C5

Target species:
Cattle

Pig
Rabbit
Equid
Sheep
Goat
Poultry

Route of administration:
Oral use

Product details

Active substance and strength:
TARAXACUM OFFICINALE D3
0.13 gram(s) / 1.00 millilitre(s)

BERBERIS VULGARIS C3
0.13 gram(s) / 1.00 millilitre(s)

SOLIDAGO VIRGAUREA C3
0.13 gram(s) / 1.00 millilitre(s)

CHINA D3
0.13 gram(s) / 1.00 millilitre(s)

Hydrastis canadensis C3
0.13 gram(s) / 1.00 millilitre(s)

Cynara scolymus D3
0.13 gram(s) / 1.00 millilitre(s)

SILYBUM MARIANUM C3
0.13 gram(s) / 1.00 millilitre(s)

Chelidonium majus C5
0.13 gram(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
- **Equid**
  - 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:
15/06/2012

Manufacturing sites for batch release:
Boiron

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

Authorisation number:
FR/V/2245672 4/2012

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.