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ALCARUMIN

Authorised

- Calcium hydroxide
- ILLICIUM VERUM
- Rhubarb
- Magnesium oxide
- Calcium carbonate
- CINCHONA OFFICINALIS BARK
- Cobaltous carbonate basic
- Cyberlindnera jadinii (Sartory, R. Sartory, Jos. Weill & J. Mey.) Minter, whole

Product identification

Medicine name:
ALCARUMIN

Active substance:
Calcium hydroxide
ILLICIUM VERUM
Rhubarb
Magnesium oxide
Calcium carbonate
CINCHONA OFFICINALIS BARK
Cobaltous carbonate basic
Cyberlindnera jadinii (Sartory, R. Sartory, Jos. Weill & J. Mey.) Minter, whole

Target species:

Cattle
Goat
Sheep

Route of administration:
In drinking water use

Product details

Active substance and strength:
Calcium hydroxide
28.80 milligram(s) / 1.00 gram(s)

ILLICIUM VERUM
28.80 milligram(s) / 1.00 gram(s)

Rhubarb
14.40 milligram(s) / 1.00 gram(s)

Magnesium oxide
43.20 milligram(s) / 1.00 gram(s)

Calcium carbonate
247.80 milligram(s) / 1.00 gram(s)

CINCHONA OFFICINALIS BARK
28.80 milligram(s) / 1.00 gram(s)

Cobaltous carbonate basic
0.34 milligram(s) / 1.00 gram(s)

Cyberlindnera jadinii (Sartory, R. Sartory, Jos. Weill & J. Mey.) Minter, whole
288.20 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for oral suspension

Withdrawal period by route of administration:
In drinking water use:

- **Cattle**
  - Milk. 0 day

- **Cattle**
  - Meat and offal. 0 day
- **Goat**
  - Milk. 0 day
- **Goat**
  - Meat and offal. 0 day
- **Sheep**
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA02AF02

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Italy

Package description:
Available only in Italian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Azienda Terapeutica Italiana A.T.I. S.r.l.

Marketing authorisation date:
28/05/1962

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
28/05/1962

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

Documents

Combined File of all Documents

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