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Tropfen bei chronischem Nasenfluss "Mag. Doskar" für Tiere

Authorised

- HYDRARGYRUM BIIODATUM D3
- KALIUM BICHROMICUM D4
- HEPAR SULFURIS D12
- HYDRASTIS CANADENSIS D3
- ACIDUM SILICICUM D12
- SULFUR IODATUM D4
- STANNUM JODATUM D8
- THUJA OCCIDENTALIS D6

Product identification

Medicine name:
Tropfen bei chronischem Nasenfluss "Mag. Doskar" für Tiere

Active substance:
HYDRARGYRUM BIIODATUM D3

KALIUM BICHROMICUM D4

HEPAR SULFURIS D12

HYDRASTIS CANADENSIS D3

ACIDUM SILICICUM D12

SULFUR IODATUM D4

STANNUM JODATUM D8

THUJA OCCIDENTALIS D6

Target species:
Cattle
Dog
Horse
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
HYDRARGYRUM BIIODATUM D3
3.00 gram(s) / 107.00 millilitre(s)

KALIUM BICHROMICUM D4
10.00 gram(s) / 107.00 millilitre(s)

HEPAR SULFURIS D12
19.00 gram(s) / 107.00 millilitre(s)

HYDRASTIS CANADENSIS D3
1.00 gram(s) / 107.00 millilitre(s)

ACIDUM SILICICUM D12
19.00 gram(s) / 107.00 millilitre(s)

SULFUR IODATUM D4
10.00 gram(s) / 107.00 millilitre(s)

STANNUM JODATUM D8
19.00 gram(s) / 107.00 millilitre(s)

THUJA OCCIDENTALIS D6
19.00 gram(s) / 107.00 millilitre(s)

Pharmaceutical form:
Oral drops, solution

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QV03AX

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

Authorisation status:
Valid

Authorised in:
Austria

Package description:
Available only in German

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:
Doskar e.U.

Marketing authorisation date:
7/12/1999

Manufacturing sites for batch release:
Doskar e.U.

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-30072

Date of authorisation status change:
7/12/1999

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

Documents

Package Leaflet

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Summary of Product Characteristics

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

Labelling

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