Sascupreel T ad us. vet. Tablette Hund, Katze

Authorised

- ACONITUM NAPELLUS D6
- AMANITA MUSCARIA D4
- AMMONIUM BROMATUM D4
- ATROPINUM SULFURICUM D6
- CITRULLUS COLOCYNTHIS D4
- CUPRUM SULFURICUM D6
- GELSEMIUM SEMPERVIRENS D6
- MAGNESIUM PHOSPHORICUM D6
- MATRICARIA RECUTITA D4
- PASSIFLORA INCARNATA D4
- VERATRUM ALBUM D6

Product identification

Medicine name:
Sascupreel T ad us. vet. Tablette Hund, Katze

Active substance:
ACONITUM NAPELLUS D6

AMANITA MUSCARIA D4

AMMONIUM BROMATUM D4

ATROPINUM SULFURICUM D6

CITRULLUS COLOCYNTHIS D4

CUPRUM SULFURICUM D6

GELSEMIUM SEMPERVIRENS D6

MAGNESIUM PHOSPHORICUM D6

MATRICARIA RECUTITA D4

PASSIFLORA INCARNATA D4

VERATRUM ALBUM D6

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
ACONITUM NAPELLUS D6
56.00 milligram(s) / 1.00 Tablet

AMANITA MUSCARIA D4
14.00 milligram(s) / 1.00 Tablet

AMMONIUM BROMATUM D4
28.00 milligram(s) / 1.00 Tablet

ATROPINUM SULFURICUM D6
28.00 milligram(s) / 1.00 Tablet

CITRULLUS COLOCYNTHIS D4
28.00 milligram(s) / 1.00 Tablet

CUPRUM SULFURICUM D6
14.00 milligram(s) / 1.00 Tablet

GELSEMIUM SEMPERVIRENS D6
28.00 milligram(s) / 1.00 Tablet

MAGNESIUM PHOSPHORICUM D6
28.00 milligram(s) / 1.00 Tablet

MATRICARIA RECUTITA D4
14.00 milligram(s) / 1.00 Tablet

PASSIFLORA INCARNATA D4
28.00 milligram(s) / 1.00 Tablet

VERATRUM ALBUM D6
28.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

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

Authorisation status:
Valid

Authorised in:
Germany

Package description:
Available only in German
Available only in German
Available only in German

Additional information

Entitlement type:
Homeopathic Registration

Marketing authorisation holder:
Biologische Heilmittel Heel GmbH

Marketing authorisation date:
25/09/2023

Manufacturing sites for batch release:
Biologische Heilmittel Heel GmbH

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
V7012300.00.00

Date of authorisation status change:
25/09/2023

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

Documents

Package Leaflet

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