

Rhus/Aconitum comp. PlantaVet Flüssige Verdünnung zur Injektion für Pferde, Rinder, Schweine, Schafe, Ziegen

Authorised

- RHUS TOXICODENDRON D5
- MANDRAGORA OFFICINARUM E RADICE FERM 34D DIL. D5 (HAB, VS. 34D)
- LEONTOPODIUM ALPINUM E PLANTA TOTA FERM 36 DIL. D3 (HAB, VS. 36)
- GRANIT DIL. D9 (HAB, VS. 6)
- GELSEMIUM SEMPERVIRENS E RHIZOMA FERM 35B DIL. D3 (HAB, VS. 35B)
- ACONITUM NAPELLUS E TUBERE FERM 33C DIL. D5 (HAB, VS. 33C)

Product identification

Medicine name:

Rhus/Aconitum comp. PlantaVet Flüssige Verdünnung zur Injektion für Pferde, Rinder, Schweine, Schafe, Ziegen

Active substance:

RHUS TOXICODENDRON D5

MANDRAGORA OFFICINARUM E RADICE FERM 34D DIL. D5 (HAB, VS. 34D)

LEONTOPODIUM ALPINUM E PLANTA TOTA FERM 36 DIL. D3 (HAB, VS. 36)

GRANIT DIL. D9 (HAB, VS. 6)

GELSEMIUM SEMPERVIRENS E RHIZOMA FERM 35B DIL. D3 (HAB, VS. 35B)

ACONITUM NAPELLUS E TUBERE FERM 33C DIL. D5 (HAB, VS. 33C)

Target species:

Cattle

Goat

Sheep

Horse

Pig

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

RHUS TOXICODENDRON D5

1.00 gram(s) / 10.00 millilitre(s)

MANDRAGORA OFFICINARUM E RADICE FERM 34D DIL. D5 (HAB, VS. 34D)

1.00 gram(s) / 10.00 millilitre(s)

LEONTOPODIUM ALPINUM E PLANTA TOTA FERM 36 DIL. D3 (HAB, VS. 36)

1.00 gram(s) / 10.00 millilitre(s)

GRANIT DIL. D9 (HAB, VS. 6)

1.00 gram(s) / 10.00 millilitre(s)

GELSEMIUM SEMPERVIRENS E RHIZOMA FERM 35B DIL. D3 (HAB, VS. 35B)

1.00 gram(s) / 10.00 millilitre(s)

ACONITUM NAPELLUS E TUBERE FERM 33C DIL. D5 (HAB, VS. 33C)

1.00 gram(s) / 10.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Germany

Package description:

Available only in German

Additional information

Entitlement type:

Homeopathic Registration

Marketing authorisation holder:

SaluVet GmbH

Marketing authorisation date:

27/11/2020

Manufacturing sites for batch release:

Wala-Heilmittel GmbH

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

402712.00.00

Date of authorisation status change:

27/11/2020

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

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