File downloaded on 2026-11-07
Source URL: https://medicines.health.europa.eu/veterinary/en/600000100836

ZEEL, injekcinis tirpalas

Authorised

- CARTILAGO SUIS D6
- FUNICULUS UMBILICALIS SUIS D6
- EMBRYO SUIS
- PLACENTA SUIS
- SYMPHYTUM OFFICINALE D6
- SOLANUM DULCAMARA D3
- NADIDUM D8
- COENZYM A D8
- SANGUINARIA CANADENSIS D4
- ARNICA MONTANA
- SULFUR D6
- NATRIUM DIETHYLOXALACETICUM D8
- ACIDUM ALPHA-KETOGLUTARICUM D8
- RHUS TOXICODENDRON D2

Product identification

Medicine name:
ZEEL, injekcinis tirpalas

Active substance:
CARTILAGO SUIS D6

FUNICULUS UMBILICALIS SUIS D6

EMBRYO SUIS

PLACENTA SUIS

SYMPHYTUM OFFICINALE D6

SOLANUM DULCAMARA D3

NADIDUM D8

COENZYM A D8

SANGUINARIA CANADENSIS D4

ARNICA MONTANA

SULFUR D6

NATRIUM DIETHYLOXALACETICUM D8

ACIDUM ALPHA-KETOGLUTARICUM D8

RHUS TOXICODENDRON D2

Target species:
Horse
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
CARTILAGO SUIS D6
5.00 microlitre(s) / 1.00 Ampoule

FUNICULUS UMBILICALIS SUIS D6
5.00 microlitre(s) / 1.00 Ampoule

EMBRYO SUIS
5.00 microlitre(s) / 1.00 Ampoule

PLACENTA SUIS
5.00 microlitre(s) / 1.00 Ampoule

SYMPHYTUM OFFICINALE D6
25.00 microlitre(s) / 1.00 Ampoule

SOLANUM DULCAMARA D3
25.00 microlitre(s) / 1.00 Ampoule

NADIDUM D8
5.00 microlitre(s) / 1.00 Ampoule

COENZYM A D8
5.00 microlitre(s) / 1.00 Ampoule

SANGUINARIA CANADENSIS D4
7.50 microlitre(s) / 1.00 Ampoule

ARNICA MONTANA
50.00 microlitre(s) / 1.00 Ampoule

SULFUR D6
9.00 microlitre(s) / 1.00 Ampoule

NATRIUM DIETHYLOXALACETICUM D8
5.00 microlitre(s) / 1.00 Ampoule

ACIDUM ALPHA-KETOGLUTARICUM D8
5.00 microlitre(s) / 1.00 Ampoule

RHUS TOXICODENDRON D2
25.00 microlitre(s) / 1.00 Ampoule

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QV03

Legal status of supply:
This information is not available for this product.

Authorisation status:
Valid

Authorised in:
Lithuania

Available in:
Lithuania

Package description:
Available only in Lithuanian
Available only in Lithuanian

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:
Biologische Heilmittel Heel GmbH

Marketing authorisation date:
23/03/2003

Manufacturing sites for batch release:
Biologische Heilmittel Heel GmbH

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/03/1539/001-002

Date of authorisation status change:
30/03/2026

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

Documents

Combined File of all Documents

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