Ginseng-logoplex injekció

Authorised

- STROPHANTHUS GRATUS D3
- Symphytum officinale D1
- RUBIA TINCTORUM D6
- RHUS TOXICODENDRON D2
- Gelsemium sempervirens D1
- SOLANUM DULCAMARA D2
- CONIUM MACULATUM D2
- ANAMIRTA COCCULUS D2
- Bryonia D1
- ATROPA BELLA-DONNA D2
- ARNICA MONTANA D1
- ACONITUM NAPELLUS D2
- PANAX QUINQUEFOLIUS

Product identification

Medicine name:
Ginseng-logoplex injekció

Active substance:
STROPHANTHUS GRATUS D3

Symphytum officinale D1

RUBIA TINCTORUM D6

RHUS TOXICODENDRON D2

Gelsemium sempervirens D1

SOLANUM DULCAMARA D2

CONIUM MACULATUM D2

ANAMIRTA COCCULUS D2

Bryonia D1

ATROPA BELLA-DONNA D2

ARNICA MONTANA D1

ACONITUM NAPELLUS D2

PANAX QUINQUEFOLIUS

Target species:

Cattle
Sheep
Goat
Horse
Pig
Dog
Cat

Route of administration:

Intramuscular use

Product details

Active substance and strength:

STROPHANTHUS GRATUS D3
0.50 milligram(s) / 1.00 millilitre(s)

Symphytum officinale D1
2.70 milligram(s) / 1.00 millilitre(s)

RUBIA TINCTORUM D6
2.70 milligram(s) / 1.00 millilitre(s)

RHUS TOXICODENDRON D2
2.70 milligram(s) / 1.00 millilitre(s)

Gelsemium sempervirens D1
2.70 milligram(s) / 1.00 millilitre(s)

SOLANUM DULCAMARA D2
2.70 milligram(s) / 1.00 millilitre(s)

CONIUM MACULATUM D2
2.70 milligram(s) / 1.00 millilitre(s)

ANAMIRTA COCCULUS D2
2.70 milligram(s) / 1.00 millilitre(s)

Bryonia D1
2.70 milligram(s) / 1.00 millilitre(s)

ATROPA BELLA-DONNA D2
2.70 milligram(s) / 1.00 millilitre(s)

ARNICA MONTANA D1
2.70 milligram(s) / 1.00 millilitre(s)

ACONITUM NAPELLUS D2
2.70 milligram(s) / 1.00 millilitre(s)

PANAX QUINQUEFOLIUS
2.70 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Hungary

Package description:
Available only in Hungarian
Available only in Hungarian

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:
Medicus Partner Kft.

Marketing authorisation date:
8/12/1997

Manufacturing sites for batch release:
Cefak KG

Responsible authority:
Directorate Of Veterinary Medicinal Products

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

Date of authorisation status change:
8/12/1997

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