

# Phytolacca-logoplex injekció

Authorised

- ACIDUM SILICICUM D10
- ECHINACEA D4
- CALCIUM SULFURICUM D12
- BRYONIA CRETICA D8
- ATROPA BELLA-DONNA D6
- ARNICA MONTANA D6
- ACIDUM FORMICICUM D4
- Phytolacca americana D5

## Product identification

**Medicine name:**

Phytolacca-logoplex injekció

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**Active substance:**

ACIDUM SILICICUM D10

ECHINACEA D4

CALCIUM SULFURICUM D12

BRYONIA CRETICA D8

ATROPA BELLA-DONNA D6

ARNICA MONTANA D6

ACIDUM FORMICICUM D4

Phytolacca americana D5

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**Target species:**

Cattle

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**Route of administration:**

Intramuscular use

Subcutaneous use

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**Product details****Active substance and strength:**

ACIDUM SILICICUM D10

100.00 milligram(s) / 1.00 millilitre(s)

ECHINACEA D4

100.00 milligram(s) / 1.00 millilitre(s)

CALCIUM SULFURICUM D12

100.00 milligram(s) / 1.00 millilitre(s)

BRYONIA CRETICA D8

100.00 milligram(s) / 1.00 millilitre(s)

ATROPA BELLA-DONNA D6

100.00 milligram(s) / 1.00 millilitre(s)

ARNICA MONTANA D6

100.00 milligram(s) / 1.00 millilitre(s)

ACIDUM FORMICICUM D4

100.00 milligram(s) / 1.00 millilitre(s)

Phytolacca americana D5

100.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Intramuscular use:**

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**Cattle**

- Meat and offal. 0 day

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Hungary

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**Package description:**

Available only in Hungarian

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Homeopathic medicinal products (Article 19 of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Medicus Partner Kft.

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**Marketing authorisation date:**

8/12/1997

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**Manufacturing sites for batch release:**

TheraSelect GmbH

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**Responsible authority:**

Directorate Of Veterinary Medicinal Products

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

8/12/1997

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)