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PHOSPHOR-HOMACCORD, injekcinis tirpalas

Authorised

- PARIS QUADRIFOLIA D200
- PARIS QUADRIFOLIA D30
- PARIS QUADRIFOLIA D10
- PARIS QUADRIFOLIA D6
- ARGENTUM NITRICUM D200
- ARGENTUM NITRICUM D30
- ARGENTUM NITRICUM D10
- PHOSPHORUS D200
- PHOSPHORUS D30
- PHOSPHORUS D10

Product identification

Medicine name:
PHOSPHOR-HOMACCORD, injekcinis tirpalas

Active substance:
PARIS QUADRIFOLIA D200

PARIS QUADRIFOLIA D30

PARIS QUADRIFOLIA D10

PARIS QUADRIFOLIA D6

ARGENTUM NITRICUM D200

ARGENTUM NITRICUM D30

ARGENTUM NITRICUM D10

PHOSPHORUS D200

PHOSPHORUS D30

PHOSPHORUS D10

Target species:
Horse
Cattle
Pig
Sheep
Goat
Dog
Cat

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
PARIS QUADRIFOLIA D200
0.02 millilitre(s) / 1.00 Ampoule

PARIS QUADRIFOLIA D30
0.02 millilitre(s) / 1.00 Ampoule

PARIS QUADRIFOLIA D10
0.02 millilitre(s) / 1.00 Ampoule

PARIS QUADRIFOLIA D6
0.02 millilitre(s) / 1.00 Ampoule

ARGENTUM NITRICUM D200
0.02 millilitre(s) / 1.00 Ampoule

ARGENTUM NITRICUM D30
0.02 millilitre(s) / 1.00 Ampoule

ARGENTUM NITRICUM D10
0.02 millilitre(s) / 1.00 Ampoule

PHOSPHORUS D200
0.02 millilitre(s) / 1.00 Ampoule

PHOSPHORUS D30
0.02 millilitre(s) / 1.00 Ampoule

PHOSPHORUS D10
0.02 millilitre(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/2/03/1538/001-002

Date of authorisation status change:
25/03/2008

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

Documents

Summary of Product Characteristics

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