

Depedin suspensija injekcijām zirgiem, liellopiem, cūkām, suņiem un kaķiem

Authorised

- Prednisolone acetate
- Dexamethasone

Product identification

Medicine name:

Depedin suspensija injekcijām zirgiem, liellopiem, cūkām, suņiem un kaķiem

Active substance:

Prednisolone acetate

Dexamethasone

Target species:

Horse

Cattle

Pig

Dog

Cat

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Prednisolone acetate

7.50 milligram(s) / 1.00 millilitre(s)

Dexamethasone

2.50 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

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Horse

- Meat and offal. 8 day
- Milk. no withdrawal period

Nelietot ķēvēm, kuru pienu paredzēts izmantot cilvēku uzturā.

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Cattle

- Meat and offal. 8 day
- Milk. 24 hour

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Pig

- Meat and offal. 6 day
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB30

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Latvia

Package description:

Available only in [Latvian](#)

Available only in [Latvian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:

Veyx Pharma GmbH

Marketing authorisation date:

9/09/1992

Manufacturing sites for batch release:

Veyx Pharma GmbH

Veyx-Pharma B.V.

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/NRP/92/0216

Date of authorisation status change:

9/09/1992

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

Documents

Labelling

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Summary of Product Characteristics

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

Package Leaflet

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