

TETANUSAN 50 % oldatos injekció A.U.V.

Authorised

- Magnesium chloride hexahydrate
- Calcium gluconate
- Sodium glycerophosphate pentahydrate

Product identification

Medicine name:

TETANUSAN 50 % oldatos injekció A.U.V.

Active substance:

Magnesium chloride hexahydrate

Calcium gluconate

Sodium glycerophosphate pentahydrate

Target species:

Cattle

Horse

Route of administration:

Intravenous use

Product details

Active substance and strength:

Magnesium chloride hexahydrate
125.00 milligram(s) / 1.00 millilitre(s)

Calcium gluconate
440.00 milligram(s) / 1.00 millilitre(s)

Sodium glycerophosphate pentahydrate
20.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intravenous use:

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Cattle

- Milk. 0 day
- Meat and offal. 0 day

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Horse

- Meat and offal. 0 day
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12CX99

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Hungary

Package description:

Available only in Hungarian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Kon-Pharma GmbH

Marketing authorisation date:

5/06/1996

Manufacturing sites for batch release:

Bela-Pharm GmbH & Co. KG

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

5/06/1996

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

Documents

Summary of Product Characteristics

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Package Leaflet

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