

Fysiologische Zoutoplossing Dechra Oplossing voor injectie

Authorised

- Sodium chloride

Product identification

Medicine name:

Fysiologische Zoutoplossing Dechra Oplossing voor injectie

Active substance:

Sodium chloride

Target species:

Horse (foal)

Cattle (calf)

Cat

Cattle

Horse

Pig

Dog

Route of administration:

Intravenous use

Subcutaneous use

Product details

Active substance and strength:

Sodium chloride

9.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intravenous use:

-

Horse (foal)

- Meat and offal. 0 day

-

Cattle (calf)

- Meat and offal. 0 day

-

Cattle

- Meat and offal. 0 day 0 days

- Milk. 0 day 0 days

-

Horse

- Meat and offal. 0 day 0 days

- Milk. 0 day 0 days

-

Pig

- Meat and offal. 0 day 0 days

Subcutaneous use:

-

Cattle (calf)

- Meat and offal. 0 day

-

Horse (foal)

- Meat and offal. 0 day

-

Cattle

- Meat and offal. 0 day 0 days

- Milk. 0 day 0 days

-

Pig

- Meat and offal. 0 day 0 days

-

Horse

- Meat and offal. 0 day 0 days

- Milk. 0 day 0 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QB05XA03

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Available in:

Belgium

Package description:

Fysiologische Zoutoplossing Dechra 1 Vial with 10 l of Solution for injection

Fysiologische Zoutoplossing Dechra 1 Vial with 1 l of Solution for injection

Fysiologische Zoutoplossing Dechra 1 Vial with 750 ml of Solution for injection

Fysiologische Zoutoplossing Dechra 1 Vial with 500 ml of Solution for injection

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Dechra Veterinary Products

Marketing authorisation date:

30/10/2006

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V288294

Date of authorisation status change:

20/04/2016

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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