

Sodu Chlorek 0,9% WET Baxter 9 g/1000 ml Roztwór do infuzji

Authorised

- Sodium chloride

Product identification

Medicine name:

Sodu Chlorek 0,9% WET Baxter 9 g/1000 ml Roztwór do infuzji

Active substance:

Sodium chloride

Target species:

Goat
Horse
Cat
Pig
Cattle
Dog
Sheep

Route of administration:

Intraperitoneal use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

Sodium chloride

9.00 gram(s) / 1000.00 millilitre(s)

Pharmaceutical form:

Solution for infusion

Withdrawal period by route of administration:

Intraperitoneal use:

-

Goat

- Meat and offal. 0 day

-

Horse

- Meat and offal. 0 day

-

Pig

- Meat and offal. 0 day

-

Cattle

- Meat and offal. 0 day

-

Sheep

- Meat and offal. 0 day

Intravenous use:

-

Goat

- Meat and offal. 0 day

-

Horse

- Meat and offal. 0 day

•

Pig

- Meat and offal. 0 day

•

Cattle

- Meat and offal. 0 day

•

Sheep

- Meat and offal. 0 day

Subcutaneous use:

•

Goat

- Meat and offal. 0 day

•

Horse

- Meat and offal. 0 day

•

Pig

- Meat and offal. 0 day

•

Cattle

- Meat and offal. 0 day

•

Sheep

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QB05BB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Package description:

Available only in Polish

Available only in Polish

Available only in Polish

Available only in Polish

Available only in Polish

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:

Baxter Polska Sp. z o.o.

Marketing authorisation date:

12/08/1996

Manufacturing sites for batch release:

Bieffe Medital S.A.

Bieffe Medital S.p.A.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

0278

Date of authorisation status change:

12/08/1996

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

Documents

Labelling

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

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.