

Vetivex 1 (9 mg/ml) solution for infusion for cattle, horses, dogs and cats

Authorised

- Sodium chloride

Product identification

Medicine name:

Vetivex 1 (9 mg/ml) solution for infusion for cattle, horses, dogs and cats

Active substance:

Sodium chloride

Target species:

Cattle

Dog

Horse

Cat

Route of administration:

Intravenous use

Product details

Active substance and strength:

Sodium chloride

9.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for infusion

Withdrawal period by route of administration:

Intravenous use:

-

Cattle

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

-

Horse

- 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:

Sweden

Package description:

Polyvinylchloride infusion bag overwrapped with polypropylene.Pack sizes: boxes containing 40 x 100 ml

Polyvinylchloride infusion bag overwrapped with polypropylene.Pack sizes: boxes containing 15 x 500 ml

Polyvinylchloride infusion bag overwrapped with polypropylene.Pack sizes: boxes containing 5 x 2000 ml.

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Polyvinylchloride infusion bag overwrapped with polypropylene.Pack sizes: boxes containing 5 x 2000 ml.

Polyvinylchloride infusion bag overwrapped with polypropylene.Pack sizes: boxes containing 12 x 1000 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Dechra Regulatory B.V.

Marketing authorisation date:

17/10/2013

Manufacturing sites for batch release:

Industria Farmaceutica Galenica Senese S.r.l.

Infomed Fluids S.R.L.

Responsible authority:

Swedish Medical Products Agency

Authorisation number:

48070

Date of authorisation status change:

17/10/2013

Reference member state:

Ireland

Procedure number:

IE/V/0511/001

Concerned member states:

Belgium Denmark France Germany Netherlands Sweden

United Kingdom (Northern Ireland)

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

Documents

Package Leaflet

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Summary of Product Characteristics

English (PDF)

Published on: 7/06/2026

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Labelling

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Combined File of all Documents