

Vetivex 11 (Hartmann's) solution for infusion for cattle, horses, dogs and cats

Authorised

- Sodium lactate
- Sodium chloride
- Potassium chloride
- Calcium chloride dihydrate

Product identification

Medicine name:

Vetivex 11 (Hartmann's) solution for infusion for cattle, horses, dogs and cats

Active substance:

Sodium lactate

Sodium chloride

Potassium chloride

Calcium chloride dihydrate

Target species:

Cattle

Dog

Horse

Cat

Route of administration:

Intravenous use

Product details

Active substance and strength:

Sodium lactate

3.20 milligram(s) / 1.00 millilitre(s)

Sodium chloride

6.00 milligram(s) / 1.00 millilitre(s)

Potassium chloride

0.40 milligram(s) / 1.00 millilitre(s)

Calcium chloride dihydrate

0.27 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:

Germany

Package description:

Polyvinylchloride infusion bag overwrapped with polypropylene. All pack sizes have two ports. Pack size: Individual fluid bag of 5000 ml, each supplied with a package leaflet

Polyvinylchloride infusion bags overwrapped with polypropylene. All pack sizes have two ports. Pack sizes: boxes containing 2 x 5000 ml.

Polyvinylchloride infusion bags overwrapped with polypropylene. All pack sizes have two ports. Pack sizes: boxes containing 15 x 500 ml

Polyvinylchloride infusion bags overwrapped with polypropylene. All pack sizes have two ports. Pack sizes: boxes containing 3 x 3000 ml

Polyvinylchloride infusion bags overwrapped with polypropylene. All pack sizes have two ports. In place of the additive port on the 5000 ml combipack is a combi port. This enables two such bags to be connected in sequence and volumes greater than 5000 ml to be administered during one infusion. Pack sizes: boxes containing 2 x 5000 ml combi.

Polyvinylchloride infusion bags overwrapped with polypropylene. All pack sizes have two ports. Pack sizes: boxes containing 4 x 3000 ml.

Polyvinylchloride infusion bags overwrapped with polypropylene. All pack sizes have two ports. Pack sizes: boxes containing 10 x 1000 ml

Polyvinylchloride infusion bags overwrapped with polypropylene. All pack sizes have two ports. Pack sizes: boxes containing 20 x 500 ml

Polyvinylchloride infusion bags overwrapped with polypropylene. All pack sizes have two ports. Pack sizes: boxes containing 20 x 250 ml

Polyvinylchloride infusion bag overwrapped with polypropylene. All pack sizes have two ports. In place of the additive port on the 5000 ml combipack is a combi port. This enables two such bag to be connected in sequence and volumes greater than 5000 ml to be administered during one infusion. Pack size: 5000 ml combi, each supplied with a package leaflet

Polyvinylchloride infusion bag overwrapped with polypropylene. All pack sizes have two ports. Pack size: Individual fluid bag of 3000 ml, each supplied with a package leaflet

Polyvinylchloride infusion bag overwrapped with polypropylene. All pack sizes have two ports. Pack size: Individual fluid bag of 1000 ml, each supplied with a package leaflet

Polyvinylchloride infusion bag overwrapped with polypropylene.All pack sizes have two ports.Pack size: Individual fluid bag of 500 ml, each supplied with a package leaflet

Polyvinylchloride infusion bag overwrapped with polypropylene.All pack sizes have two ports. Pack size: Individual fluid bag of 250 ml, each supplied with a package leaflet.

Polyvinylchloride infusion bags overwrapped with polypropylene.All pack sizes have two ports.Pack sizes: boxes containing 30 x 250 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:

11/06/2013

Manufacturing sites for batch release:

Industria Farmaceutica Galenica Senese S.r.l.

Infomed Fluids S.R.L.

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

401806.00.00

Date of authorisation status change:

20/09/2018

Reference member state:

Ireland

Procedure number:

IE/V/0512/001

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

Combined File of all Documents

English (PDF)

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Summary of Product Characteristics