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

Authorised

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

Product identification

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

Active substance:
Calcium chloride dihydrate
Potassium chloride
Sodium chloride
Sodium lactate

Target species:
Cattle
Dog
Horse
Cat

Route of administration:
Intravenous use

Product details

Active substance and strength:

Calcium chloride dihydrate
0.27 milligram(s) / 1.00 millilitre(s)

Potassium chloride
0.40 milligram(s) / 1.00 millilitre(s)

Sodium chloride
6.00 milligram(s) / 1.00 millilitre(s)

Sodium lactate
3.20 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
- **Dog**
- **Horse**
  - Meat and offal. 0 day
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QB05BB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

Package description:

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 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 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 3000 ml, each supplied with a package leaflet

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 bags overwrapped with polypropylene.All pack sizes have two ports.Pack sizes: boxes containing 20 x 250 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 10 x 1000 ml

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. 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 3 x 3000 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 2 x 5000 ml.
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

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 Limited

Marketing authorisation date:
20/06/2013

Manufacturing sites for batch release:
Industria Farmaceutica Galenica Senese S.r.l.
Infomed Fluids S.R.L.

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
10434/4080

Date of authorisation status change:
20/06/2013

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

Summary of Product Characteristics