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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:
Sweden

Available in:
Sweden

Package description:
Available only in Swedish
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Polyvinylchloride infusion bags overwrapped with polypropylene.All pack sizes have two ports.Pack sizes: boxes containing 30 x 250 ml
Polyvinylchloride infusion bags overwrapped with polypropylene.All pack sizes have two ports.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:
Infomed Fluids S.R.L.
Industria Farmaceutica Galenica Senese S.r.l.

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
48074

Date of authorisation status change:
17/10/2013

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

English (PDF)
Published on: 7/06/2026

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Package Leaflet

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Combined File of all Documents

Labelling

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