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
Infusolec Infusionsvätska, lösning

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

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Authorisation status:

Valid

Authorised in:

Sweden

Package description:
Available only in Swedish
Available only in Swedish
Available only in Swedish
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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:
This information is not available for this product.

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:

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: 3/05/2024
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Package Leaflet

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