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UISCE-JECT, 100 % v/v, solvent for parenteral use

Authorised

- Water for injections

Product identification

Medicine name:
UISCE-JECT, 100 % v/v, solvent for parenteral use

Active substance:
Water for injections

Target species:
Cattle
Dog
Sheep
Horse
Cat
Pig

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Water for injections
100.00 millilitre(s) / 100.00 millilitre(s)

Pharmaceutical form:
Solvent for parenteral use

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Pig**
  - Meat and offal. 0 day

Intravenous use:

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

- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Pig**
  - Meat and offal. 0 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QV07AB

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Ireland

Package description:
100 ml polypropylene vials with grey bromobutyl rubber stoppers and aluminium flip-off seals. Pack sizes: Box with 10 vials of 100 ml
100 ml polypropylene vials with grey bromobutyl rubber stoppers and aluminium flip-off seals. Pack sizes: Box with 1 vial of 100 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:
Duggan Veterinary Supplies Limited

Marketing authorisation date:
15/09/2017

Manufacturing sites for batch release:
S P Veterinaria S.A.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10400/003/001

Date of authorisation status change:

15/09/2017

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

Documents

Summary of Product Characteristics