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T-61 SOLUCION INYECTABLE PARA INDUCCION DE LA EUTANASIA EN ANIMALES

Authorised

- Mebezonium iodide
- Embutramide
- Tetracaine hydrochloride

Product identification

Medicine name:
T-61 SOLUCION INYECTABLE PARA INDUCCION DE LA EUTANASIA EN ANIMALES

Active substance:
Mebezonium iodide
Embutramide
Tetracaine hydrochloride

Target species:
Horse
Dog
Cat
Ruminant
Ornamental bird
Fur animals
Laboratory animals

Route of administration:

Intravenous use
Intracardiac use
Intrapulmonary use

Product details

Active substance and strength:
Mebezonium iodide
50.00 milligram(s)/millilitre / 1.00 millilitre(s)

Embutramide
200.00 milligram(s)/millilitre / 1.00 millilitre(s)

Tetracaine hydrochloride
5.00 milligram(s)/millilitre / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intravenous use:

- **Horse**
  - Meat and offal. no withdrawal period

Carne: Se tomarán las medidas adecuadas para garantizar que los cuerpos de los animales muertos tratados con este medicamento veterinario y los subproductos de estos animales no entren en la cadena alimentaria y no se utilicen para el consumo humano o ani

- **Ruminant**
  - Meat and offal. no withdrawal period

Carne: Se tomarán las medidas adecuadas para garantizar que los cuerpos de los animales muertos tratados con este medicamento veterinario y los subproductos de estos animales no entren en la cadena alimentaria y no se utilicen para el consumo humano o ani

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN51AX50

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

Authorisation status:
Valid

Authorised in:
Spain

Available in:
Spain

Package description:
Available only in Spanish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Merck Sharp & Dohme Animal Health S.L.

Marketing authorisation date:
7/02/1994

Manufacturing sites for batch release:
Intervet International GmbH

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
873 ESP

Date of authorisation status change:
7/02/1994

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

Documents

Summary of Product Characteristics

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Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.