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

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