

# 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

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**Active substance:**

Mebezonium iodide

Embutramide

Tetracaine hydrochloride

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**Target species:**

Horse

Dog

Cat

Ruminant

Ornamental bird

Fur animals

Laboratory animals

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**Route of administration:**

Intravenous use  
Intracardiac use  
Intrapulmonary use

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## 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)

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### **Pharmaceutical form:**

Solution for injection

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### **Withdrawal period by route of administration:**

#### **Intravenous use:**

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##### **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

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##### **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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QN51AX50

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Spain

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**Available in:**

Spain

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**Package description:**

Available only in Spanish

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Merck Sharp & Dohme Animal Health S.L.

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**Marketing authorisation date:**

7/02/1994

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**Manufacturing sites for batch release:**

Intervet International GmbH

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**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

873 ESP

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**Date of authorisation status change:**

7/02/1994

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Summary of Product Characteristics

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

### 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.