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# T 61, injekcinis tirpalas

Authorised

- Embutramide
- Mebezonium iodide
- Tetracaine hydrochloride

## Product identification

**Medicine name:**
T 61, injekcinis tirpalas

**Active substance:**
Embutramide
Mebezonium iodide
Tetracaine hydrochloride

**Target species:**
Dog
Large animals
Cat
Birds

**Route of administration:**
Intravenous use
Intrapulmonary use

## Product details

**Active substance and strength:**
Embutramide
200.00 milligram(s) / 1.00 millilitre(s)

Mebezonium iodide
50.00 milligram(s) / 1.00 millilitre(s)

Tetracaine hydrochloride
5.00 milligram(s) / 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:**

- **Dog**
  - All relevant tissues. no withdrawal period

T 61 euthanized animals must be disposed of in accordance with national requirements.

- **Large animals**
  - All relevant tissues. no withdrawal period

Do not use in animals intended for human consumption. T 61 euthanized animals must be disposed of in accordance with national requirements.

**Intrapulmonary use:**

- **Dog**
  - All relevant tissues. no withdrawal period

T 61 euthanized animals must be disposed of in accordance with national requirements.

- **Cat**
  - All relevant tissues. no withdrawal period

T 61 euthanized animals must be disposed of in accordance with national requirements.

- **Birds**
  - All relevant tissues. no withdrawal period

Do not use in animals intended for human consumption. T 61 euthanized animals must be disposed of in accordance with national requirements.

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QN51AX50

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

**Authorisation status:**
Valid

**Authorised in:**
Lithuania

**Package description:**
Available only in Lithuanian

## Additional information

**Entitlement type:**
Marketing Authorisation

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

**Marketing authorisation holder:**
Intervet International B.V.

**Marketing authorisation date:**
15/04/2002

**Manufacturing sites for batch release:**
Intervet International GmbH

**Responsible authority:**
State Food And Veterinary Service

**Authorisation number:**
This information is not available for this product.

**Date of authorisation status change:**
15/04/2007

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

## Documents

RV1399.pdf