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)

Pharmaceutical form:
Solution for injection

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