

EUTHAPHEN™

(Euthanasia Solution)

- For use in dogs for humane, painless and rapid euthanasia.
- A non-sterile solution containing pentobarbital sodium and phenytoin sodium as the active ingredients.
- Each mL contains 390 mg pentobarbital sodium and 50 mg phenytoin sodium.
- Rhodamine B, a bluish-red fluorescent dye, is included in the formulation to help distinguish it from parenteral drugs intended for therapeutic use.
- Available in 100 mL multiple-dose vials.



Dosage and Administration

- Dosage: Dogs: 1 mL for each 10 pounds of body weight
- Administration: Intravenous injection is preferred. Intracardiac injection may be made when intravenous injection is impractical.

To order, please contact your Dechra representative or call (866) 683-0660.

For more information, please visit www.dechra-us.com

Veterinary Technical Support available (866) 933-2472

Nonurgent Technical Support available via email support@dechra.com

IMPORTANT SAFETY INFORMATION

For canine euthanasia only. Must not be used for therapeutic purposes. Do not use in animals intended for food. Keep out of reach of children. If eye contact, flush eyes with water and seek medical attention. Caution should be exercised to avoid contact of the drug with open wounds or accidental self-inflicted injections. Euthanasia may sometimes be delayed in dogs with severe cardiac or circulatory deficiencies.

ENVIRONMENTAL HAZARD: This product is toxic to wildlife. Birds and mammals feeding on treated animals may be killed. Euthanized animals must be properly disposed of by deep burial, incineration, or other method in compliance with state and local laws, to prevent consumption of carcass material by scavenging wildlife.

Refer to the prescribing information for complete details or visit www.dechra-us.com

EUTHAPHEN™

(Euthanasia Solution)



FOR DOGS ONLY

CAUTION: Federal law restricts this drug to use by or on the order of a licensed veterinarian.

DESCRIPTION:

A non-sterile solution containing pentobarbital sodium and phenytoin sodium as the active ingredients. Rhodamine B, a bluish-red fluorescent dye, is included in the formulation to help distinguish it from parenteral drugs intended for therapeutic use. Although the solution is not sterile, benzyl alcohol, a bacteriostat, is included to retard the growth of microorganisms.

Each mL contains: *Active ingredients:* 390 mg pentobarbital sodium (barbituric acid derivative), 50 mg phenytoin sodium. *Inactive ingredients:* 10% ethyl alcohol, 18% propylene glycol, 0.003688 mg rhodamine B, 2% benzyl alcohol (preservative), purified water q.s. Sodium hydroxide and/or hydrochloric acid may be added to adjust pH.

ACTIONS:

EUTHAPHEN contains two active ingredients which are chemically compatible but pharmacologically different. Each ingredient acts in such a manner so as to cause humane, painless and rapid euthanasia. Euthanasia is due to cerebral death in conjunction with respiratory arrest and circulatory collapse. Cerebral death occurs prior to cessation of cardiac activity.

When administered intravenously, pentobarbital sodium produces rapid anesthetic action. There is a smooth and rapid onset of unconsciousness. At the lethal dose, there is depression of vital medullary respiratory and vasomotor centers.

When administered intravenously, phenytoin sodium produces toxic signs of cardiovascular collapse and/or central nervous system depression. Hypotension occurs when the drug is administered rapidly.

PHARMACODYNAMIC ACTIVITY:

The sequence of events leading to humane, painless and rapid euthanasia following intravenous injection of EUTHAPHEN is similar to that following intravenous injection of pentobarbital sodium or other barbituric derivatives. Within seconds, unconsciousness is induced with simultaneous collapse of the dog. This stage rapidly progresses to deep anesthesia with concomitant reduction in the blood pressure. A few seconds later, breathing stops, due to depression of the medullary respiratory center; encephalographic activity becomes isoelectric, indicating cerebral death; and then cardiac activity ceases.

Phenytoin sodium exerts its effect during the deep anesthesia stage caused by the pentobarbital sodium. This ingredient, due to its cardiotoxic properties, hastens the stoppage of electrical activity in the heart.

INDICATIONS:

For use in dogs for humane, painless and rapid euthanasia.

WARNING:

For canine euthanasia only. Must not be used for therapeutic purposes. Do not use in animals intended for food.

ENVIRONMENTAL HAZARD:

This product is toxic to wildlife. Birds and mammals feeding on treated animals may be killed. Euthanized animals must be properly disposed of by deep burial, incineration, or other method in compliance with state and local laws, to prevent consumption of carcass material by scavenging wildlife.

HUMAN WARNING:

Caution should be exercised to avoid contact of the drug with open wounds or accidental self-inflicted injections. Keep out of reach of children. If eye contact, flush eyes with water and seek medical attention.

PRECAUTIONS:

Euthanasia may sometimes be delayed in dogs with severe cardiac or circulatory deficiencies. This may be explained by the impaired movement of the drug to its site of action. An occasional dog may elicit reflex responses manifested by motor movement; however, an unconscious animal does not experience pain, because the cerebral cortex is not functioning.

When restraint may cause the dog pain, injury or anxiety, or danger to the person making the injection, prior use of tranquilizing or immobilizing drugs may be necessary.

DOSAGE AND ADMINISTRATION:

Dosage: Dogs: 1 mL for each 10 pounds of body weight.

Administration: Intravenous injection is preferred. Intracardiac injection may be made when intravenous injection is impractical, as in a very small dog, or in a comatose dog with impaired vascular functions. Good injection skill is necessary for intracardiac injection.

The calculated dose should be given in a single bolus injection.

For intravenous injection, a needle of sufficient gauge to ensure intravenous placement of the entire dose should be used.

The use of a Luer-Lok® syringe is recommended to prevent accidental exposure to needle/syringe separation.

To report suspected adverse drug events, for technical assistance or to obtain a copy of the Safety Data Sheet (SDS), contact Dechra at (866) 933-2472.

For additional information about adverse drug experience reporting for animal drugs, contact FDA at 1-888-FDA-VETS or online at <http://www.fda.gov/reportanimalae>

HOW SUPPLIED: EUTHAPHEN is available in 100 mL multiple-dose vials.

NDC 17033-088-10

Manufactured by a non-sterilizing process.

STORAGE: Store between 15° and 30°C (59°F and 86°F).

Manufactured for:

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Approved by FDA under ANADA # 200-280

Luer-Lok® is a registered trademark of Becton, Dickinson and Company.

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