

EUTHAPHEN- euthanasia injection, solution
Dechra Veterinary Products, LLC

EUTHAPHEN C-III (EUTHANASIA SOLUTION)

Approved by FDA under ANADA # 200-280

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™ C-III (Euthanasia) Solution 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 solution 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 wild life. 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.

HOW SUPPLIED: EUTHAPHEN[™] Solution is available in 100 mL multiple-dose vials.

Manufactured by a non-sterilizing process.

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

Manufactured for:

DECHRA VETERINARY PRODUCTS, LLC
Overland Park, KS 66211

Rev.: 02/2024

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

Human Warning: Refer to package insert for important human risk information.

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

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.

Manufactured for: Dechra Veterinary Products
7015 College Boulevard, Suite 525
Overland Park, KS 66211 USA

NDC 17033-088-10

EUTHAPHEN™ 
(pentobarbital sodium and phenytoin sodium)
Euthanasia Solution
FOR DOGS ONLY

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

Approved by FDA under ANADA # 200-280

Net Contents:  **100 mL** 

Manufactured by a non-sterilizing process. Multiple-Dose Vial.
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.
For Intravenous or Intracardiac Use. Read product information sheet carefully. See warnings on left panel.

Rev, November 2023 3  17033 08810 4 

EUTHAPHEN

euthanasia injection, solution

Product Information

Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:17033-088
Route of Administration	INTRAVENOUS, INTRACARDIAC	DEA Schedule	CIII

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PENTOBARBITAL SODIUM (UNII: NJJ0475N0S) (PENTOBARBITAL - UNII:I4744080IR)	PENTOBARBITAL SODIUM	390 mg in 1 mL
PHENYTOIN SODIUM (UNII: 4182431BJH) (PHENYTOIN - UNII:6158TKW0C5)	PHENYTOIN SODIUM	50 mg in 1 mL

Product Characteristics

Color	red (bluish-red)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17033-088-10	100 mL in 1 VIAL, MULTI-DOSE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANADA	ANADA200280	01/11/2023	

Labeler - Dechra Veterinary Products, LLC (362142734)

Revised: 2/2024

Dechra Veterinary Products, LLC