

TEMPLATE FOR RELEVANT LABEL PARTICULARS (RLPs) (*Veterinary Products*)

Select appropriate:

× New Product (include all applicable RLPs)



RLP
Approved



Signal heading:	PRESCRIPTION ANIMAL REMEDY KEEP OUT OF REACH OF CHILDREN FOR ANIMAL TREATMENT ONLY
Product name:	Urilin Syrup for Dogs
Active constituent/s:	50.0 mg/mL Phenylpropanolamine hydrochloride
Statement of claims:	For use in bitches to control urinary incontinence associated with acquired urethral sphincter incompetence.
Net contents:	45 mL 100 mL
Directions for Use Heading:	To be used by, or under the direction of, a registered veterinarian.
Restrictions:	-
Contraindications:	This product is contraindicated for use in animals treated with non-selective monoamine oxidase inhibitors. This product is contraindicated in animals with known hypersensitivity to the active substance.
Precautions:	Because phenylpropanolamine is a sympathomimetic agent, it may affect the cardiovascular system, especially blood pressure and heart rate, and therefore should be used with caution in animals with cardiovascular diseases. Care should be exercised in treating animals with severe renal or hepatic insufficiency, diabetes mellitus, hyperadrenocorticism, glaucoma or other metabolic disorders. In bitches less than 1 year old the possibility of anatomical disorders contributing to incontinence should be considered prior to treatment. It is not appropriate to use the product for the behavioural cause of inappropriate urination Care should be exercised in administering the product with other sympathomimetic drugs, anticholinergic drugs, tricyclic antidepressants or specific type B monoamine oxidase inhibitors. DO NOT USE in pregnant or lactating bitches
Side effects:	Adverse reactions: If you notice any serious effects or other effects not mentioned in this leaflet, please inform your veterinary surgeon. In some dogs, loose stools, liquid diarrhoea, a decrease in appetite, arrhythmia and collapse have been reported following treatment with phenylpropanolamine. Occasional nausea and vomiting have also been reported. As phenylpropanolamine is a sympathomimetic drug it is possible to produce a wide range of effects, most of which mimic the results of excess stimulation of the sympathetic nervous system (e.g. effects on the heart rate and blood pressure). Dizziness and restlessness have been reported in some dogs following treatment. Hypersensitivity may occur in very rare cases. Overdose An overdose could produce signs of excessive stimulation of the sympathetic nervous system.

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	Treatment should be symptomatic. Alpha-adrenergic blockers may be appropriate in the case of severe overdose. However, no specific recommendation on drugs or dosages can be given.
Dosage & administration:	For oral administration only. The recommended dose is 1mg phenylpropanolamine hydrochloride /kg body weight three times daily in the feed, corresponding to 0.1 ml Urilin Syrup/5 kg body weight three times daily. 1 drop for every 2.34 kg body weight three times daily in feed. Discard unused portion three months after first use.
General directions:	Do not shake the bottle, simply invert it over the food and count the required number of drops. The efficacy of phenylpropanolamine has only been demonstrated in ovariectomised bitches.
Withholding Period/s:	Not applicable
Trade Advice:	Not applicable
Safety Directions:	
First Aid:	If poisoning occurs, contact a doctor or Poisons Information Centre, phone Australia 131126.
Additional user safety:	Special precautions for the person administering the veterinary medicinal product to animals. Phenylpropanolamine hydrochloride is toxic when overdoses are ingested. Adverse effects may include dizziness, headache, nausea, insomnia or restlessness, and increased blood pressure. High overdose may be fatal, especially in children. In the event of accidental skin contact, wash the contaminated area with soap and water. Wash hands after use of the product. In the event of accidental eye contact, rinse the eye with clean water for about 15 minutes and seek medical advice. To avoid accidental ingestion the product must be used and kept out of the reach and sight of children. Always replace the cap firmly after use to ensure that the child resistant closure operates correctly. In the event of accidental ingestion, seek immediate medical attention showing the doctor the package leaflet
Environmental statements:	Not applicable
Disposal:	Dispose of empty containers by wrapping with paper and putting in garbage
Storage:	Store below 25°C (Air Conditioning). Protect from light by keeping the container in the outer carton. Do not freeze. Discard unused portion three months after first use.
Name & address:	

The following is for APVMA use only:

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