

**PARTICULARS TO APPEAR ON THE OUTER PACKAGE – CARTON 100 ml
bottle and 30 ml bottle**

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

IncoVet Syrup, 40 mg/ml, dogs

2. STATEMENT OF ACTIVE SUBSTANCES

Each ml contains:

Active substance:

Phenylpropanolamine (as hydrochloride)40.28 mg/ml
(Equivalent to 50 mg phenylpropanolamine hydrochloride)

3. PACKAGE SIZE

30 ml

100 ml

4. TARGET SPECIES

Dogs



5. INDICATIONS

6. ROUTES OF ADMINISTRATION

Oral administration.

7. WITHDRAWAL PERIODS

8. EXPIRY DATE

Exp. {mm/yyyy}

Once opened, use within 3 months

Once opened, use by: / /

9. SPECIAL STORAGE PRECAUTIONS

Do not store above 25°C.

Keep the bottle in outer carton in order to protect from light.

10. THE WORDS “READ THE PACKAGE LEAFLET BEFORE USE”

Read the package leaflet before use.

11. THE WORDS “FOR ANIMAL TREATMENT ONLY”

For animal treatment only.

12. THE WORDS “KEEP OUT OF THE SIGHT AND REACH OF CHILDREN”

Keep out of the sight and reach of children.

13. NAME OF THE MARKETING AUTHORISATION HOLDER

VETOQUINOL UK Limited

14. MARKETING AUTHORISATION NUMBER

Vm 08007/5001

15. BATCH NUMBER

Lot {number}

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE – LABEL 100ml bottle

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

IncoVet Syrup, 40 mg/ml, dogs

2. STATEMENT OF ACTIVE SUBSTANCES

Each ml contains:

Active substance:

Phenylpropanolamine (as hydrochloride)40.28 mg/ml
(Equivalent to 50 mg phenylpropanolamine hydrochloride)

3. TARGET SPECIES

Dogs



4. ROUTES OF ADMINISTRATION

Oral administration.

Read the package leaflet before use.

5. WITHDRAWAL PERIODS

6. EXPIRY DATE

Exp. {mm/yyyy}

Once opened, use within 3 months.

Once opened, use by: / /

7. SPECIAL STORAGE PRECAUTIONS

Do not store above 25°C.

Keep the bottle in outer carton in order to protect from light

8. NAME OF THE MARKETING AUTHORISATION HOLDER

VETOQUINOL UK Limited

9. BATCH NUMBER

Lot {number}

MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING
UNITS – LABEL 30ml bottle

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

IncoVet Syrup, 40 mg/ml, dogs

Phenylpropanolamine



2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES

Each ml contains:

Active substance:

Phenylpropanolamine (as hydrochloride).....40.28 mg/ml

(Equivalent to 50 mg phenylpropanolamine hydrochloride)

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp. {mm/yyyy}Once opened, use within 3 months.

Once opened, use by: / /

PARTICULARS TO APPEAR ON THE PACKAGE LEAFLET:

PACKAGE LEAFLET

1. Name of the veterinary medicinal product

IncoVet Syrup, 40 mg/ml, dogs

2. Composition

Each ml contains:

Active substance:

Phenylpropanolamine 40.28 mg
(Equivalent to 50 mg phenylpropanolamine hydrochloride)

Colourless to slightly yellow-brown solution.

3. Target species

Dogs.



4. Indications for use

Treatment of urinary incontinence associated with urethral sphincter incompetence in the bitch.

Efficacy has only been demonstrated in ovariohysterectomised bitches.

5. Contraindications

The use of the veterinary medicinal product is not appropriate for the treatment of behavioural causes of inappropriate urination. Do not use in patients treated with non-selective monoamine oxidase inhibitors.

Do not use in cases of hypersensitivity to the active substance or to any of the excipients.

6. Special warnings

Special warnings:

For Animal Treatment Only

Special precautions for safe use in the target species:

Due to the very low doses to be administered, and to avoid any risk of overdose, the animal must be weighed, and the recommended doses must be respected.

Phenylpropanolamine, a sympathomimetic drug, may affect the cardiovascular system, especially blood pressure and heart rate, and 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, hyperthyroidism 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.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:

Phenylpropanolamine hydrochloride is toxic when ingested in higher doses. Adverse effects may include dizziness, headache, nausea, insomnia or restlessness, and increased blood pressure.

Accidental ingestion by a child may be fatal.

To avoid accidental ingestion, the veterinary medicinal product must be used and kept out of the sight and reach of children. Always replace the cap securely after use and store the syringe and bottle inside the cardboard box at all times.

In case of accidental ingestion, seek medical attention immediately and show the package leaflet or the label to the physician.

In case of accidental skin contact, wash the contaminated area with soap and water. Wash hands after handling the veterinary medicinal product.

This veterinary medicinal product may cause eye irritation.

In case of accidental eye contact, rinse the eye with clean water for about 15 minutes and seek medical advice immediately and show the package leaflet or the label to the physician.

Pregnancy and lactation:

The safety of the veterinary medicinal product has not been established during pregnancy or lactation.

Pregnancy and lactation:

Do not use during pregnancy or lactation.

Interaction with other medicinal products and other forms of interaction:

Care should be exercised in administering the veterinary medicinal product with other sympathomimetic drugs, anticholinergic drugs, tricyclic antidepressants or specific type B monoamine oxidase inhibitors.

Overdose:

In healthy dogs, no side effects were observed at up to 5 times the recommended dosage. However, an overdose of phenylpropanolamine could produce symptoms of excessive stimulation of the sympathetic nervous system. 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.

7. Adverse events

Dogs:

<p>Rare (1 to 10 animals / 10,000 animals treated):</p>	<p>Diarrhoea¹, loose stool, emesis (vomiting) Lethargy</p>
<p>Very rare (<1 animal / 10,000 animals treated, including isolated reports):</p>	<p>Tachycardia², high blood pressure² (hypertension²), arrhythmia (irregular heartbeat) Proteinuria² (presence of protein in the urine) Wobbliness, ataxia (difficulty coordinating movements), seizure Decreased appetite Collapse, polydipsia (increased thirst) Aggression, hyperactivity (agitation) Hypersensitivity Polyuria (Increased urination)</p>

¹ Liquid

² Effects of sympathomimetics producing excessive stimulation of the sympathetic nervous system

Reporting adverse events is important. It allows continuous safety monitoring of a product. If you notice any side effects, even those not already listed in this package leaflet, or you think that the medicine has not worked, please contact, in the first instance, your veterinarian. You can also report any adverse events to the marketing authorisation holder using the contact details at the end of this leaflet, or via your national reporting system at:

Website: <https://www.gov.uk/report-veterinary-medicine-problem/animal-reacts-medicine>

e-mail: adverse.events@vmd.gov.uk

8. Dosage for each species, routes and method of administration

Oral use.

To ensure a correct dosage, body weight should be determined as accurately as possible.

The recommended dose for the veterinary medicinal product is 1mg/kg bodyweight 3 times daily in the feed, corresponding to 0.1 ml of the veterinary medicinal product / 5 kg bodyweight (i.e., a graduation of the provided syringe for 5 kg), 3 times daily.

The absorption rate is increased if the veterinary medicinal product is administered to fasted dogs.

9. Advice on correct administration

None.

Instructions for use



1. Remove the childproof safety cap by pushing down firmly and turning anti-clockwise.



2. Take the dosage syringe, with plunger pushed inwards, and insert the end into the opening of the stopper. Push down firmly.



3. Hold at an angle, do not fully invert the bottle, draw IncoVet Syrup into the syringe slowly to avoid air bubbles. Stop at the relevant mark on the plunger for the volume of IncoVet Syrup required.



4. Turn bottle the right way up and take hold of the lower part of the syringe, close to the neck of the bottle. Carefully remove the syringe from the bottle using a turning action.



5. Hold the syringe over your dog's food and push the plunger inwards as far as it will go to ensure you use the full dose of IncoVet Syrup required.

6. Replace the cap onto the bottle and turn clockwise to tighten. Store bottle safely, at room temperature, out of the sight and reach of children.

7. Wipe the end of the syringe with a clean tissue or paper towel and keep in a clean place. If the syringe needs cleaning, remove the plunger and rinse both parts with warm water.

8. Dry thoroughly, ensuring that the inside of the syringe is dry before replacing the plunger.

10. Withdrawal periods

Not applicable.

11. Special storage precautions

Keep out of the sight and reach of children.

Do not store above 25°C.

Keep the bottle in outer carton in order to protect from light.

Do not use this veterinary medicinal product after the expiry date which is stated on the label or on the carton after EXP. The expiry date refers to the last day of that month.

Shelf life after first opening the container: 3 months.

When the container is breached/opened for the first time, using the in-use shelf-life which is specified on this package leaflet, the date on which any product remaining in the container should be discarded should be determined. This discard date should be written in the space provided.

12. Special precautions for disposal

Medicines should not be disposed of via wastewater.

Use take-back schemes for the disposal of any unused veterinary medicinal product or waste materials derived thereof in accordance with local requirements and with any applicable national collection systems. These measures should help to protect the environment.

Ask your veterinary surgeon how to dispose of medicines no longer required.

13. Classification of veterinary medicinal products

Veterinary medicinal product subject to prescription.

14. MARKETING AUTHORISATION NUMBER AND PACK SIZES

Vm 08007/5001

Box with 1 bottle of 30 ml with a syringe of 1.5 ml

Box with 1 bottle of 100 ml with a syringe of 1.5 ml

Not all pack sizes may be marketed.

15. PID LINK (Do not print heading)

[The following statement must be included where reference to the European Union Product Database is included on the product information. This statement is relevant to both UK(GB) and UK(NI) products:]

Find more product information by searching for the 'Product Information Database' on www.gov.uk.

16. Contact details

Marketing authorisation holder and contact details to report suspected adverse reactions:

Vetoquinol UK Limited
Steadings Barn
Pury Hill Business Park
Nr Alderton
Towcester
Northamptonshire
NN12 7LS

Manufacturer responsible for batch release:

VETOQUINOL S.A.
Magny-Vernois
F-70200 Lure
FRANCE

17. Other information

POM-V

Pharmacodynamic properties

The clinical effect of phenylpropanolamine in urinary incontinence is based on its stimulation effect on α -adrenergic receptors. This causes an increase in, and a stabilisation of, the closure pressure in the urethra, which is innervated mainly by adrenergic nerves.

Phenylpropanolamine is a racemic mixture of D and L enantiomers

Pharmacokinetic particulars

In the dog, the mean half-life of Phenylpropanolamine is approximately 3 hours with maximal plasma concentrations being found after approximately 1 hour. No accumulation of phenylpropanolamine has been observed after a dose of 1 mg/kg 3 times daily over 15 days.

When the product is administered to a fasted dog, bioavailability is increased significantly.

Gavin Hall
Approved: 27 March 2026