

PRODUCT MONOGRAPH

PrLUPIN-TOLVAPTAN
tolvaptan tablets

Tablets, 15 mg, 30 mg, 45 mg, 60 mg and 90 mg

Vasopressin V₂-receptor Antagonist

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LUPIN-TOLVAPTAN
tolvaptan tablets

PART I: HEALTH PROFESSIONAL INFORMATION

SUMMARY PRODUCT INFORMATION

Route of Administration	Dosage Form / Strength	Non-medicinal Ingredients
Oral	Tablets, 15 mg, 30 mg, 45 mg, 60 mg and 90 mg	Corn starch, hydroxypropyl cellulose, lactose monohydrate, low-substituted hydroxypropyl cellulose, magnesium stearate, microcrystalline cellulose, and FD&C Blue No. 2 Aluminum Lake as colorant

INDICATIONS AND CLINICAL USE

LUPIN-TOLVAPTAN (tolvaptan) is indicated to slow the progression of kidney enlargement and kidney function decline in patients with autosomal dominant polycystic kidney disease (ADPKD). In ADPKD, kidney enlargement reflects renal cyst burden.

In order to select patients who might best benefit from the effects of tolvaptan, clinical trials evaluated ADPKD patients having a total kidney volume (TKV) ≥ 750 mL, and/or renal function corresponding to a CKD-EPI eGFR ≥ 25 mL/min/1.73m², at the time of initiation of treatment.

LUPIN-TOLVAPTAN treatment should be initiated and monitored under the supervision of a nephrologist or specialist with expertise in the management of patients with ADPKD and a full understanding of the benefits and risks of tolvaptan therapy including hepatic toxicity and monitoring requirements.

Careful consideration and discussion of the appropriateness of LUPIN-TOLVAPTAN treatment should be undertaken between the prescriber and patient before initiation of therapy, taking into account the potential benefits and risks of treatment. Upon mutual agreement to undertake treatment with LUPIN-TOLVAPTAN, a signed, duly- documented, manufacturer and product-specific patient-prescriber agreement (PPAF) is required outlining relevant patient selection criteria to be considered, expected benefits and risks of treatment, and the need for mandatory hepatic function monitoring (see [WARNINGS AND PRECAUTIONS, Hepatotoxicity](#), and [DOSAGE AND ADMINISTRATION](#)).

LUPIN-TOLVAPTAN Controlled Hepatic Safety Monitoring and Distribution Programme

LUPIN-TOLVAPTAN is available for treatment of patients with ADPKD only through a manufacturer and product-specific controlled hepatic safety monitoring and distribution (HSMD) programme conducted and maintained by, or for, the market authorisation holder of LUPIN-TOLVAPTAN. A duly signed manufacturer and product-specific PPAF is required for enrollment in the HSMD programme. For more information on the programme, please call 1-

866-488-6017.

Geriatrics (> 65 years of age): The safety and effectiveness of tolvaptan in geriatric patients has not been established.

Pediatrics (< 18 years of age): Tolvaptan has not been studied in pediatric patients with ADPKD. Its use is not recommended in this patient population.

CONTRAINDICATIONS

LUPIN-TOLVAPTAN is contraindicated in:

- Patients who have been asked to permanently discontinue tolvaptan in the past
- Patients with known or suspected hypersensitivity to tolvaptan, benzazepine or benzazepine derivatives (e.g. mirtazapine) or to any of the excipients. For a complete listing, see [DOSAGE FORMS, COMPOSITION AND PACKAGING](#).
- Patients with hypovolemia
- Patients with hypernatremia
- Patients with anuria (see [WARNINGS AND PRECAUTIONS, Renal Impairment](#))
- Patients who do not have access to fluids or who cannot respond to the physiologic sensation of thirst
- Patients with a history, signs or symptoms of significant liver impairment or injury, excluding uncomplicated polycystic liver disease (see [WARNINGS AND PRECAUTIONS, Hepatotoxicity](#), and/or [Hepatic Impairment](#), and [DOSAGE AND ADMINISTRATION, Dosing Considerations, Hepatic Impairment](#))
- Concomitant use of strong CYP3A inhibitors, e.g., ketoconazole, itraconazole, clarithromycin, telithromycin, ritonavir, indinavir, nelfinavir, saquinavir, nefazodone (see [DRUG INTERACTIONS, Drug-Drug Interactions](#))
- Pregnancy (See [WARNINGS AND PRECAUTIONS, Special Populations, Pregnant Women](#))
- Nursing women (See [WARNINGS AND PRECAUTIONS, Special Populations, Nursing Women](#))
- Patients with one of the following rare hereditary diseases: Galactose intolerance, Lapp lactase deficiency or Glucose-galactose malabsorption because lactose is a non-medicinal ingredient in LUPIN-TOLVAPTAN.

WARNING: IDIOSYNCRATIC HEPATIC TOXICITY

Tolvaptan use has led to idiosyncratic elevations of blood alanine and aspartate aminotransferases (ALT & AST), rarely associated with concomitant elevations of total

bilirubin (see [Hepatotoxicity](#), below). To help mitigate the risk of liver injury, blood testing for hepatic transaminases and total bilirubin is required prior to initiation of LUPIN-TOLVAPTAN, then hepatic transaminases continuing monthly for 18 months, every 3 months for the next 12 months, and then every 3-6 months thereafter during treatment with LUPIN-TOLVAPTAN (see [DOSAGE AND ADMINISTRATION, Dosing Considerations](#); See [WARNINGS AND PRECAUTIONS, Hepatotoxicity](#)). Therefore, LUPIN-TOLVAPTAN is available for treatment of patients with ADPKD only through a controlled hepatic safety monitoring and distribution (HSMD) programme conducted and maintained by, or for, the market authorization holder of LUPIN-TOLVAPTAN.

WARNINGS AND PRECAUTIONS

Dehydration

Due to a prominent aquaretic effect, treatment with tolvaptan may result in dehydration which constitutes a risk factor for renal dysfunction. If dehydration becomes evident, take appropriate action which may include the need to interrupt or reduce the dose of tolvaptan and increase fluid intake. Special care should be taken in patients having diseases that impair appropriate fluid intake or who are at an increased risk of water loss, e.g. in case of vomiting or diarrhea. LUPIN-TOLVAPTAN should not be prescribed to patients who cannot perceive or respond to thirst (see [CONTRAINDICATIONS](#)).

LUPIN-TOLVAPTAN may cause undesirable effects related to water loss such as thirst, polyuria, nocturia, and pollakiuria. Therefore, it is imperative that patients should have access to water (or other aqueous fluids) and be able to drink sufficient amounts of these fluids. Patients should be instructed to drink water or other aqueous fluids at the first sign of thirst in order to avoid excessive thirst or dehydration (see [DOSAGE AND ADMINISTRATION](#)).

Patients should be generally encouraged to drink water while taking LUPIN-TOLVAPTAN to avoid development of dehydration or hypernatremia, and to improve tolerability of tolvaptan.

Drugs metabolised by CYP-3A or transported by P-gp

Tolvaptan is a substrate of CYP-3A and co-administration with CYP-3A inhibitors or CYP-3A inducers may lead to a change in exposure. Patient response should be monitored and the dose adjusted accordingly, as appropriate.

CYP-3A Inhibitors

CYP3A4 is the predominant enzyme involved in tolvaptan metabolism. Concomitant use with strong CYP3A4 inhibitors is contraindicated as this may lead to a significant increase in tolvaptan exposure.

In an interaction trial, co-administration of ketoconazole, a strong inhibitor of CYP3A4, significantly inhibited the metabolism of tolvaptan, increasing mean tolvaptan exposure up to 5.4-fold (as determined by AUC_{∞}). (see [CONTRAINDICATIONS](#)).

Concomitant use with moderate CYP-3A inhibitors, i.e., verapamil, fluconazole and erythromycin, requires lowered dosing of LUPIN-TOLVAPTAN (see [DRUG INTERACTIONS](#), and [DOSAGE AND ADMINISTRATION, CYP-3A Inhibitors](#)).

LUPIN-TOLVAPTAN should not be taken with grapefruit juice.

CYP-3A Inducers

Concomitant use of LUPIN-TOLVAPTAN with strong CYP-3A inducers should be avoided, e.g. rifampin, phenytoin, carbamazepine, St. John's Wort (see [DRUG INTERACTIONS](#), and [DOSAGE AND ADMINISTRATION, CYP-3A Inducers](#)).

P-gp Inhibitors

Reduction in the dose of LUPIN-TOLVAPTAN may be required in patients concomitantly treated with P-glycoprotein (P-gp) inhibitors, such as cyclosporine and quinidine, based on clinical response (see [DRUG INTERACTIONS](#), and [DOSAGE AND ADMINISTRATION, P-gp Inhibitors](#)). However, concomitant use with those P-gp inhibitors that also act as strong CYP-3A inhibitors (e.g., ketoconazole, clarithromycin, ritonavir, saquinavir), is contraindicated (see [CONTRAINDICATIONS](#)).

Hepatotoxicity

Tolvaptan has been associated with idiosyncratic drug-induced hepatocellular injury, as seen by elevations of serum alanine and aspartate aminotransferases (ALT and AST), rarely associated with concomitant elevations of total bilirubin (BT).

In a double-blind, placebo-controlled trial called TEMPO 3:4 completed in patients with ADPKD, elevation (>3 x upper limit of normal [ULN]) of ALT was observed in 4.4% (42/958) of patients on tolvaptan, and 1.0% (5/484) of patients on placebo, while elevation (>3 x ULN) of AST was observed in 3.1% (30/958) of patients on tolvaptan and 0.8% (4/484) patients on placebo when monitoring for liver enzymes elevation was every 3-4 months (also, see [ADVERSE REACTIONS, Hepatic Injury](#)). Two (2/957, 0.2%) of these tolvaptan treated-patients, as well as a third patient from an extension open-label trial, exhibited increases in hepatic enzymes (>3 x ULN) with concomitant elevations in total bilirubin (>2 x ULN). The period of onset of hepatocellular injury, as reflected by ALT elevations >3 x ULN, was within 3 to 14 months after initiating treatment, and these increases were reversible, with ALT returning to <3 x ULN within 1 to 4 months. While these concomitant elevations were gradually reversible with prompt discontinuation of tolvaptan, they represent a potential for significant liver injury. Similar changes with other drugs have been associated with the potential to cause irreversible and potentially life-threatening liver injury. The incidence of hepatotoxicity does not appear to be dose related. To date, there is no evidence of hepatocellular injury with tolvaptan when used in patients not treated for ADPKD.

In post-marketing experience with tolvaptan in ADPKD, acute liver failure requiring liver transplantation has been reported.

In another double-blind placebo-controlled trial (called REPRISE) including patients with later stages of ADPKD, all patients were monitored monthly for liver enzymes elevation. In the single-blind tolvaptan run-in phase, elevations (>3 x ULN) of ALT and AST were observed in 0.2% (3/1,478) and 0.2% (3/1,477) of patients on tolvaptan, respectively. Relative to the first double-blind, placebo-controlled trial, similar elevations (>3 x ULN) of ALT (5.6% (38/681) of patients on tolvaptan and 1.2% (8/685) of patients on placebo) and AST (3.5% (24/681) of patients on tolvaptan and 0.9% (6/685) of patients on placebo) were observed in the double-blind phase. Overall, all patients recovered and no subject met Hy's laboratory criteria relative to laboratory values of potential clinical relevance, i.e. increases in hepatic enzymes (>3 x ULN)

with concomitant elevations in total bilirubin (>2 x ULN) (see [ADVERSE REACTIONS, Hepatic Injury](#)). Data suggest that monthly liver function monitoring during treatment helps detect liver enzymes elevation early on.

To mitigate the risk of significant and/or irreversible liver injury, blood testing for hepatic transaminases and total bilirubin is required prior to initiation of LUPIN-TOLVAPTAN, then hepatic transaminases continuing monthly for 18 months, every 3 months for the next 12 months, and then at 3-6 month intervals thereafter during treatment with LUPIN-TOLVAPTAN (see [DOSAGE AND ADMINISTRATION, Dosing Considerations](#)). Concurrent monitoring for symptoms that may indicate liver injury (such as fatigue, anorexia, nausea, right upper abdominal discomfort, vomiting, fever, rash, pruritus, icterus, dark urine or jaundice) is also warranted.

At the onset of symptoms or signs consistent with hepatic injury, or if abnormal ALT or AST increases are detected, LUPIN-TOLVAPTAN administration must be immediately interrupted and repeat liver tests, i.e., ALT, AST, total bilirubin, alkaline phosphatase, must be obtained as soon as possible, ideally within 48-72 hours (see [DOSAGE AND ADMINISTRATION, Dosing Considerations](#)). Testing should continue at an increased frequency until symptoms/signs/laboratory abnormalities stabilise or resolve, at which point cautious re-initiation of LUPIN-TOLVAPTAN may be considered.

Permanent discontinuation from receiving tolvaptan is a contraindication and so once a patient has been permanently discontinued from receiving tolvaptan, treatment **must never be** restarted. The permanent discontinuation status of patients should be verified prior to initiation with tolvaptan.

Anaphylaxis

In post-marketing experience, anaphylaxis (including anaphylactic shock and generalised rash) has been reported very rarely following administration of tolvaptan. This type of reaction occurred after the first administration of tolvaptan. Patients have to be carefully monitored during treatment. If an anaphylactic reaction or other serious allergic reactions occur, administration of LUPIN-TOLVAPTAN must be discontinued immediately and appropriate therapy initiated. Since hypersensitivity is a contraindication, treatment **must never be** restarted after an anaphylactic reaction or other serious allergic reactions.

Patients with known hypersensitivity reactions to benzazepines or benzazepine derivatives (e.g. benazepril, conivaptan, fenoldopam mesylate or mirtazapine) may be at risk for hypersensitivity reaction to tolvaptan. If a hypersensitivity reaction is suspected, tolvaptan should be discontinued. (see [CONTRAINDICATIONS](#)).

Hypernatremia

During treatment initiation, patients must be frequently monitored for serum sodium and volume status. If serum sodium increases above the normal range, tolvaptan treatment must be down-titrated or discontinued promptly, with serum sodium carefully monitored and appropriate clinical measures taken, as necessary (see [ADVERSE REACTIONS, Increases in Serum Sodium](#)). Concomitant use with hypertonic saline solutions or drugs that may increase serum sodium concentrations should be avoided.

Hyperkalemia

Treatment with tolvaptan is associated with an acute reduction of the extracellular fluid volume which could result in increased serum potassium. Serum potassium levels should be monitored carefully after initiation of tolvaptan, especially in those who are receiving drugs known to increase serum potassium levels, e.g. spironolactone.

Hyperuricemia

Treatment with tolvaptan may lead to increases in serum uric acid and clinical gout (see [ADVERSE REACTIONS, Increases in Serum Uric Acid](#), and [ACTION AND CLINICAL PHARMACOLOGY, Pharmacodynamics](#)). Uric acid concentrations should be evaluated prior to initiation of LUPIN-TOLVAPTAN therapy, and as indicated during treatment (see [DOSAGE AND ADMINISTRATION, Increases in Serum Uric Acid](#)).

Hypotension-related adverse events

In patients taking anti-hypertensive agents concomitantly with tolvaptan, an increased incidence of hypotension-related adverse events was observed (see [ADVERSE REACTIONS, Co-administration with Anti-Hypertensive Medications](#)), including dizziness and syncope. These findings were not seen in patients not taking anti-hypertensive drugs.

Serum sodium abnormalities

Serum sodium abnormalities, i.e., hyponatremia or hypernatremia, must be corrected prior to initiation of LUPIN-TOLVAPTAN therapy (also, see [CONTRAINDICATIONS](#)).

Vasopressin analogues

In addition to V₂-receptor mediated renal aquaretic effects, tolvaptan blocks vascular vasopressin V₂-receptors involved in the release of coagulation factors (e.g., von Willebrand factor) from endothelial cells. Therefore, the effect of vasopressin analogues such as desmopressin may be attenuated in patients using these therapies concomitantly with tolvaptan. Accordingly, it is not recommended to administer LUPIN-TOLVAPTAN together with vasopressin analogues.

Special Populations

Pregnant Women:

There are no adequate, well-controlled trials of tolvaptan in pregnant women. In animal trials with tolvaptan, cleft palate, brachymelia, microphthalmia, skeletal malformations, decreased fetal weight, delayed fetal ossification, and embryo-fetal death occurred. LUPIN-TOLVAPTAN use is contraindicated in pregnant women.

Women of childbearing potential must have effective contraceptive measures in place prior to and during LUPIN-TOLVAPTAN use.

Nursing Women:

It is not known whether tolvaptan is secreted into human milk. The presence of tolvaptan has been observed in the milk of lactating rats. Because many drugs are secreted into human milk and because of the potential for serious adverse reactions in nursing infants, treatment with LUPIN-TOLVAPTAN is contraindicated in nursing women.

Pediatrics (< 18 years of age):

Tolvaptan has not been studied in pediatric patients with ADPKD. Its use is not recommended in this patient population.

Geriatrics (> 65 years of age):

Safety and effectiveness in geriatric patients has not been studied.

Hepatic Impairment:

In hyponatremia studies, moderate or severe hepatic impairment was found to decrease the clearance and increase the volume of distribution of tolvaptan (see [CONTRAINDICATIONS](#), [DOSAGE AND ADMINISTRATION, Dosing Considerations](#), [Hepatic Impairment](#), and [ACTION AND CLINICAL PHARMACOLOGY, Hepatic Insufficiency](#)). The effects of such changes have not been studied in ADPKD patients, since these patients generally have normal liver function even in the presence of polycystic liver disease. LUPIN-TOLVAPTAN is contraindicated in patients with clinically relevant impairment of hepatic function.

Renal Impairment:

LUPIN-TOLVAPTAN is contraindicated in anuric patients, since it is not expected that tolvaptan would offer benefit in these patients. Further, no clinical trial data are available in ADPKD patients on chronic dialysis or with a creatinine clearance < 10 mL/min or with severe renal insufficiency (e.g. CKD-EPI eGFR <25 mL/min/1.73m²), including those unable to make urine (see [DOSAGE AND ADMINISTRATION, Dosing Considerations](#), [Renal Impairment](#)).

Potential for Cognitive and Motor Impairment

There are no controlled trials of the effects of tolvaptan on driving performance. When driving vehicles or using machines, it should be taken into account that occasionally dizziness, asthenia and syncope may occur.

ADVERSE REACTIONS**Adverse Drug Reaction Overview**

The adverse reaction profile of tolvaptan is mainly based on three phase III clinical trials (TEMPO 3:4, TEMPO 4:4 and REPRISE) which are briefly described in the CLINICAL TRIALS section.

In TEMPO 3:4, tolvaptan, total daily dose of 60-120 mg, has been evaluated for safety in 961 adult ADPKD patients, with approximately 2,335 patient-years of exposure to tolvaptan. A total of 836 patients were treated with tolvaptan for at least 1 year and 742 patients treated with tolvaptan had at least 3 years of exposure.

In TEMPO 4:4, an open-label extension of TEMPO 3:4, tolvaptan was evaluated for safety in 1,083 adult patients at a total daily dose of 60-120 mg, with approximately 3,281 patient-years of exposure to tolvaptan. A total of 698 and 509 patients were treated with tolvaptan for at least 1 year and at least 3 years, respectively.

In the REPRISE study, tolvaptan was evaluated for safety in 683 adult patients with approximately 612 patient-years of exposure to tolvaptan at a daily divided dose of up to 90 mg/30 mg. A total of 577 patients were treated with tolvaptan for at least 1 year.

The total clinical safety database for tolvaptan is comprised of approximately 3,226 adult patients who participated in single and multiple-dose trials in ADPKD and who had approximately 8,430 patient-years of exposure to tolvaptan.

TEMPO 3:4

In a double-blind, placebo-controlled trial known as TEMPO 3:4, the most commonly reported adverse reactions, consistent with the pharmacologic activity of tolvaptan, are thirst, polyuria, nocturia, and pollakiuria occurring in approximately 55%, 38%, 29% and 23% of patients, respectively. The most commonly reported serious adverse events (SAE) that occurred more frequently in tolvaptan patients compared to placebo patients, with $\geq 0.5\%$ difference, included increased ALT (0.9% vs. 0.4%), increased AST (0.9% vs. 0.4%), chest pain (0.8% vs. 0.4%) and headache (0.5% vs. 0%).

Adverse events that led to discontinuation of tolvaptan were reported for 15.0% of patients, compared to 4.3% of placebo-treated patients, and included polyuria, pollakiuria, nocturia, thirst, abnormal hepatic function, and fatigue.

REPRISE

In REPRISE, patients who successfully completed the single-blind tolvaptan run-in period were randomized to a double-blind phase. If an adverse event continued from the single- to double-blind period, it was considered treatment emergent in the double-blind treatment period only if the adverse event worsened, became more severe or serious, or led to discontinuation. In the single-blind tolvaptan run-in phase the most frequently reported adverse reactions were: polyuria (31.4%), thirst (28.2%), and nocturia (20.3%). In the single-blind tolvaptan run-in phase, the most frequently reported serious adverse events (SAE) were alanine aminotransferase increased (0.3%), polyuria (0.3%), and nocturia (0.2%). In the double-blind phase, the most frequently reported adverse reactions were (tolvaptan vs placebo): polyuria (5.3% vs. 1.3%), nocturia (4.7% vs. 1.6%), blood creatinine increased (4.6% vs. 4.8%), thirst (3.8% vs. 1.9%), fatigue (3.5% vs. 0.6%), headache (2.3% vs. 1.6%), ALT increased (3.2% vs. 0.9%), and AST increased (2.2% vs. 1.2%) and the most commonly reported SAE that occurred more frequently in tolvaptan patients compared to placebo patients, with $\geq 0.5\%$ difference, included hepatic enzyme increased (1.6% vs 0.1%) and alanine aminotransferase increased (1.2% vs 0.0%).

In the single-blind tolvaptan run-in phase, 6.8% of patients discontinued tolvaptan due to an adverse event which included polyuria, nocturia, thirst, fatigue and pollakiuria. In the double-blind phase, adverse events that led to discontinuation of tolvaptan were reported for 9.5% of patients, compared to 2.2% of placebo-treated patients, and included hepatic enzyme increased, ALT increased, AST increased, liver function test increased, and pollakiuria.

Idiosyncratic elevations of hepatic aminotransferases, i.e., ALT and AST, have been observed in ADPKD patients treated with tolvaptan, rarely associated with concomitant elevations in total bilirubin (see [Clinical Trial Adverse Drug Reactions](#), [Hepatic Injury](#), below, and [WARNINGS AND PRECAUTIONS, Hepatotoxicity](#)).

Clinical Trial Adverse Drug Reactions

Because clinical trials are conducted under very specific conditions the adverse reaction rates observed in the clinical trials may not reflect the rates observed in practice and should not be compared to the rates in the clinical trials of another drug. Adverse drug reaction information from clinical trials is useful for identifying drug-related adverse events and for approximating rates.

TEMPO 3:4

Table 1 below presents the incidence of treatment emergent adverse drug reactions occurring at a rate of 3% or greater and greater than placebo in TEMPO 3:4.

Table 1 Incidence of Treatment-Emergent Adverse Drug Reactions in at Least 3% of Tolvaptan Subjects and Greater than Placebo in TEMPO 3:4		
System Organ Class Preferred Term	Tolvaptan (N = 961) n (%)	Placebo (N = 483) n (%)
Cardiac disorders		
Palpitations	34 (3.5)	6 (1.2)
Gastrointestinal disorders		
Abdominal Distension	47 (4.9)	16 (3.3)
Constipation	81 (8.4)	12 (2.5)
Diarrhea	128 (13.3)	53 (11.0)
Dry Mouth	154 (16.0)	60 (12.4)
Dyspepsia	76 (7.9)	16 (3.3)
Gastroesophageal Reflux Disease	43 (4.5)	16 (3.3)
General disorders and administration site conditions		
Asthenia	57 (5.9)	27 (5.6)
Fatigue	131 (13.6)	47 (9.7)
Thirst	531 (55.3)	99 (20.5)
Metabolism and nutrition disorders		
Decreased Appetite	69 (7.2)	5 (1.0)
Hyperuricemia	37 (3.9)	9 (1.9)
Polydipsia	100 (10.4)	17 (3.5)
Musculoskeletal and connective tissue disorders		
Muscle Spasms	35 (3.6)	17 (3.5)
Nervous system disorders		
Dizziness	109 (11.3)	42 (8.7)
Headache	241 (25.1)	121 (25.1)
Psychiatric disorders		
Insomnia	55 (5.7)	21 (4.3)
Renal and urinary disorders		
Nocturia	280 (29.1)	63 (13.0)
Pollakiuria	223 (23.2)	26 (5.4)
Polyuria	368 (38.3)	83 (17.2)
Skin and subcutaneous tissue disorders		
Pruritus	33 (3.4)	13 (2.7)
Rash	40 (4.2)	9 (1.9)

REPRISE

The incidence of treatment emergent adverse drug reactions occurring at a rate of 3% or greater and greater than placebo in the REPRISE trial are presented in Table 2 below.

Table 2 Incidence of Treatment-Emergent Adverse Drug Reactions in at Least 3% of Tolvaptan Subjects and Greater than Placebo in REPRISE		
Single-blind Treatment Period (≥3% only)		
System Organ Class Preferred Term	Tolvaptan (N = 1,491) n (%)	-
Gastrointestinal Disorders		
Dry Mouth	132 (8.9)	-
General Disorders and Administration Site Conditions		
Fatigue	64 (4.3)	-
Thirst	430 (28.8)	-
Infections and Infestations		
Viral Upper Respiratory Tract Infection	47 (3.2)	-
Metabolism and Nutrition Disorders		
Polydipsia	146 (9.8)	-
Nervous System Disorders		
Headache	63 (4.2)	-
Renal and Urinary Disorders		
Nocturia	308 (20.7)	-
Pollakiuria	70 (4.7)	-
Polyuria	475 (31.9)	-
Double-blind Treatment Period (≥3% and greater than placebo)		
System Organ Class Preferred Term	Tolvaptan (N = 681) n (%)	Placebo (N = 685) n (%)
Gastrointestinal Disorders		
Abdominal Pain	25 (3.7)	15 (2.2)
Constipation	22 (3.2)	18 (2.6)
Diarrhea	47 (6.9)	23 (3.4)
General Disorders and Administration Site Conditions		
Fatigue	46 (6.8)	24 (3.5)
Thirst	27 (4.0)	13 (1.9)
Investigations		
Alanine Aminotransferase Increased	25 (3.7)	9 (1.3)
Metabolism and Nutrition Disorders		
Gout	21 (3.1)	20 (2.9)
Nervous System Disorders		
Dizziness	25 (3.7)	19 (2.8)
Renal and Urinary Disorders		
Nocturia	32 (4.7)	12 (1.8)
Polyuria	36 (5.3)	11 (1.6)

Hepatic Injury

TEMPO 3:4

Elevations (>3 x ULN) of ALT were observed in 4.4% (42/958) of patients on tolvaptan, and 1.0% (5/484) of patients on placebo in TEMPO 3:4, while elevation (>3 x ULN) of AST was observed in 3.1% (30/958) of patients on tolvaptan and 0.8% (4/484) patients on placebo when monitoring for liver enzymes elevation was every 3-4 months (see [WARNINGS AND PRECAUTIONS, Hepatotoxicity](#)). Elevations (>5 x ULN) of ALT were observed in 2.3%

(22/958) of patients on tolvaptan, and 0.4% (2/484) of placebo-treated patients, while similar elevations of AST were observed in 1.9% (18/958), and 0.4% (2/484), respectively. Elevations (>10 x ULN) of ALT were observed in 1.3% (12/958) of tolvaptan-treated patients, and (>20 x ULN) in 0.6% (6/958), with none in placebo-treated patients. Elevations (>10 x ULN) of AST were observed in 1.0% (10/958) of tolvaptan-treated patients, and (>20 x ULN) in 0.3% (3/958), with none in placebo. Two (2/957, 0.2%) of these tolvaptan treated-patients, as well as a third patient from an extension open-label trial (TEMPO 4:4), exhibited increases in hepatic enzymes (>3 x ULN) with concomitant elevations in total bilirubin (>2 x ULN). In this pivotal trial, no cases of clinically-apparent hepatic failure were reported.

REPRISE

In the single-blind tolvaptan run-in phase of REPRISE, elevations (>3 x ULN) of ALT and AST were observed in 0.2% (3/1,478) and 0.2% (3/1,477) of patients on tolvaptan, respectively. Elevation of (>5 x ULN) of ALT was observed in 0.1% (1/1,478) of patients on tolvaptan; no patients experienced ALT elevations of (>10 x ULN) or (>20 x ULN). No AST elevations of (>5 x ULN), (>10 x ULN) or (>20 x ULN) were observed.

In the double-blind phase, all patients were monitored monthly for liver enzymes elevations. Elevations (>3 x ULN) of ALT were observed in 5.6% (38/681) of patients on tolvaptan, and 1.2% (8/685) of patients on placebo in REPRISE, while elevation (>3 x ULN) of AST were observed in 3.5% (24/681) of patients on tolvaptan, and 0.9% (6/685) of patients on placebo. Elevations of (>5 x ULN) of ALT were observed in 3.4% (23/681) of patients on tolvaptan, and 0.7% (5/685) of patients on placebo, while similar elevations of AST were observed in 1.8% (12/681), and 0.6% (4/685), respectively. Elevations (>10 x ULN) of ALT or AST were observed in 1.2% (8/681) of tolvaptan-treated patients, and 0.6% (4/685) of patients on placebo. Elevations (>20 x ULN) of ALT or AST were observed in 0.1% (1/681 and 1/685, respectively) of patients on tolvaptan and placebo. For the 29 subjects in the tolvaptan group who discontinued tolvaptan after reporting a potential clinically significant increased ALT (> 3 x ULN), the values returned toward normal after tolvaptan was discontinued. For the 9 subjects in the tolvaptan group who did not discontinue the investigational medicinal product after reporting a potential clinically significant increased ALT (> 3 x ULN), the ALT values returned to normal for all except for 1 subject; that subject had 4 serious TEAEs of liver function test increased, the last of which resulted in discontinuation of tolvaptan; values returned to normal upon treatment discontinuation. Overall, no subject met the criteria for Hy's laboratory criteria relative to laboratory values of potential clinical relevance, i.e. increases in hepatic enzymes (>3 x ULN) with concomitant elevations in total bilirubin (>2 x ULN). Data suggest that monthly liver function monitoring during treatment helps detect liver enzymes elevation early on and mitigate the risk for drug- induced liver injury.

Co-administration with Anti-Hypertensive Medications

Among ADPKD patients taking anti-hypertensive medications, a higher incidence of dizziness, presyncope, and syncope was observed in patients treated with tolvaptan, compared to those treated with placebo as shown in Table 3 (also, see [WARNINGS AND PRECAUTIONS, Hypotension-related adverse events](#)).

Table 3 Incidence of Adverse Events Associated with Decreased Blood Pressure in TEMPO 3:4					
System organ class	Adverse Event	No anti-hypertensives		Anti-hypertensives	
		Tolvaptan (N=129)	Placebo (N=64)	Tolvaptan (N=832)	Placebo (N=419)
Nervous system disorders	Dizziness	8 (6.2%)	5 (7.8%)	101 (12.1%)	37 (8.8%)
	Dizziness exertional	0 (0.0%)	0 (0.0%)	1 (0.1%)	0 (0.0%)
	Dizziness postural	0 (0.0%)	0 (0.0%)	6 (0.7%)	0 (0.0%)
	Presyncope	0 (0.0%)	0 (0.0%)	3 (0.4%)	0 (0.0%)
	Syncope	3 (2.3%)	0 (0.0%)	13 (1.6%)	3 (0.7%)
Total		11 (8.5%)	5 (7.8%)	116 (13.9%)	39 (9.3%)

Adverse events are counted only once for multiple occurrences of an event.

Increases in Serum Sodium

TEMPO 3:4

The overall incidence of hypernatremia, reported as an adverse event, in TEMPO 3:4 was 2.8% in patients treated with tolvaptan, compared to 1.0% in placebo-treated patients. The incidence of serum sodium > 150 mEq/L was 4.0%, and 1.4%, respectively. None of these events led to discontinuation of tolvaptan.

Mean trough sodium levels were observed to be 1-3 mEq/L higher than placebo.

REPRISE

In the single-blind period, no incidence of hypernatremia was reported. In the double-blind treatment period of REPRISE, 0.4% of hypernatremia were reported in the tolvaptan group and 0% in the placebo group. None of these events lead to tolvaptan discontinuation.

Increases in Serum Uric Acid

TEMPO 3:4

Decreased uric acid clearance by the kidney is a known effect of tolvaptan. Gout was observed in 2.9% (20/961) of patients on tolvaptan and 1.4% (7/483) patients on placebo in TEMPO 3:4, with a higher incidence of the use of allopurinol (8.2% vs. 5.8%), benzbromarone (0.4% vs. 0.2%), and colchicine-containing medications (2.3% vs. 0.8%) to manage gout in patients on tolvaptan, compared to patients on placebo, respectively. Similarly, there was a higher incidence of increased blood uric acid (greater than 10 mg/dL), at 6.2% vs. 1.7%, reported in patients on tolvaptan compared to patients on placebo, respectively (see [DOSAGE AND ADMINISTRATION, Increases in Serum Uric Acid](#)). Mean serum uric acid levels increased about 0.9 mg/dL from baseline in patients treated with tolvaptan (see [ACTION AND CLINICAL PHARMACOLOGY, Pharmacodynamics](#)).

REPRISE

In the single-blind period, gout was reported in 0.5% (7/1,491). Gout was observed in 3.1% (21/681) of patients on tolvaptan and 2.9% (20/685) of patients on placebo in the double-blind placebo-controlled trial in ADPKD patients with late stage 2 to early stage 4 chronic kidney disease (CKD).

Less Common Clinical Trial Adverse Drug Reactions <3%

The following adverse reactions occurred in <3% of ADPKD patients treated with tolvaptan and at a rate greater than placebo in the double-blind, placebo-controlled trial (n = 961 tolvaptan; n = 483 placebo), and are not mentioned elsewhere in the labeling:

Metabolism and Nutrition Disorders: dehydration, hyperglycemia

Respiratory, Thoracic and Mediastinal Disorders: dyspnea

Post-Market Adverse Drug Reactions

Angioedema, anaphylactic shock, and generalized rash, have been reported very rarely following administration of tolvaptan for indications not related to ADPKD.

DRUG INTERACTIONS

Overview

Tolvaptan is a substrate of CYP-3A, and thus co-administration with CYP-3A inhibitors or CYP-3A inducers may lead to a change in tolvaptan exposure. Patient response should be monitored and the dose adjusted accordingly, as appropriate (see [DOSAGE AND ADMINISTRATION](#)). Tolvaptan does not inhibit or induce its own metabolism.

Non-clinical studies indicate that tolvaptan is a substrate and competitive inhibitor of p-glycoprotein; other transporters have not been studied.

There have been no trials performed to determine the potential interaction of tolvaptan with alcohol.

Drug-Drug Interactions

Effects of Other Drugs on Tolvaptan

CYP-3A Inhibitors

CYP3A4 is the predominant enzyme involved in tolvaptan metabolism. Concomitant use of tolvaptan with strong CYP3A4 inhibitors is contraindicated as this may lead to a significant increase in tolvaptan exposure. In an interaction trial, co-administration of ketoconazole, a strong inhibitor of CYP3A4, significantly inhibited the metabolism of tolvaptan, increasing mean tolvaptan exposure up to 5.4-fold (as determined by AUC_{∞}). Co-administration of a 30 mg single dose of tolvaptan with 200 mg QD ketoconazole resulted in a 440% increase in AUC and 248% increase in C_{max} for tolvaptan (see [CONTRAINDICATIONS](#)).

CYP-3A Inducers

Co-administration of a single oral dose of 240 mg of tolvaptan with 600 mg QD of rifampicin, a strong CYP-3A4 inducer, at steady state decreased tolvaptan C_{max} and AUC_t by approximately 85% (see [WARNINGS AND PRECAUTIONS, CYP-3A Inducers](#)).

P-gp Inhibitors

Co-administration of tolvaptan with P-gp inhibitors has not been studied in dedicated clinical studies (see [WARNINGS AND PRECAUTIONS, P-gp Inhibitors](#)). Concomitant use with those P-gp inhibitors that also act as strong CYP-3A inhibitors (e.g., ketoconazole, clarithromycin,

ritonavir, saquinavir), is contraindicated (see [CONTRAINDICATIONS](#))

Effects of Tolvaptan on Other Drugs

CYP-3A substrates

In healthy subjects, tolvaptan, a CYP-3A substrate, had no effect on the plasma concentrations of some other CYP-3A substrates, e.g., warfarin or amiodarone. However, tolvaptan increased plasma levels of lovastatin by 1.3- to 1.4-fold, indicating a potential effect on weak substrates of CYP-3A substrates.

Digoxin

Steady-state digoxin concentrations were statistically significantly increased (approximately 30% increase as determined by C_{max} and 20% increase as determined by AUC_{τ}), when digoxin was co-administered with multiple 60 mg doses (QD) of tolvaptan; *in vitro* studies indicate that tolvaptan is a substrate and competitive inhibitor of p-glycoprotein. Patients receiving digoxin should be evaluated for excessive digoxin effects after adding tolvaptan.

Warfarin, Amiodarone, Furosemide, and Hydrochlorothiazide

Co-administration of tolvaptan does not appear to alter the pharmacokinetics of warfarin, furosemide, hydrochlorothiazide, or amiodarone (or its active metabolite, desethylamiodarone) to a clinically significant degree.

Vasopressin Analogues

In addition to its renal aquaretic effect, tolvaptan is capable of blocking vascular vasopressin V_2 receptors involved in the release of coagulation factors (e.g., von Willebrand's factor) from endothelial cells. Therefore, the effect of vasopressin analogs such as desmopressin may be attenuated in patients using such analogs to prevent or control bleeding when co-administered with tolvaptan. Administering LUPIN-TOLVAPTAN with vasopressin analogues is not recommended.

Drug-Food Interactions

Grapefruit Juice

Co-administration of tolvaptan with 240 mL of grapefruit juice produced a doubling of peak tolvaptan concentrations (C_{max}) but had no effect on tolvaptan elimination half-life (see [WARNINGS AND PRECAUTIONS, CYP-3A Inhibitors](#)). LUPIN-TOLVAPTAN should not be taken with grapefruit juice (see [WARNINGS & PRECAUTIONS; Drugs metabolised by CYP-3A or transported by P-gp](#)).

Drug-Herb Interactions

Interactions with herbal products have not been established; however, St John's Wort should be avoided while taking LUPIN-TOLVAPTAN.

Pharmacodynamic Interactions

Concomitant Diuretic Use

Tolvaptan use alone produces a greater 24-hour urine volume than does furosemide or hydrochlorothiazide alone. However, concomitant administration of tolvaptan with furosemide or

hydrochlorothiazide results in a 24-hour volume that is similar to that after tolvaptan administration alone.

Furosemide co-administered with tolvaptan produces a similar maximal rate of urine excretion compared to furosemide alone, and 70% higher than tolvaptan alone. HCTZ co-administered with tolvaptan produces a slightly higher maximal excretion rate compared to tolvaptan alone, and 66% higher compared to HCTZ alone.

DOSAGE AND ADMINISTRATION

LUPIN-TOLVAPTAN treatment should be initiated and monitored by a physician experienced in the diagnosis and treatment of autosomal dominant polycystic kidney disease (ADPKD; see [INDICATIONS AND CLINICAL USE](#)).

Upon mutual agreement to undertake treatment with LUPIN-TOLVAPTAN, a signed, duly-documented, manufacturer and product-specific patient-prescriber agreement (PPAF) is required for all patients before initiation of LUPIN-TOLVAPTAN (see [INDICATIONS AND CLINICAL USE](#)). This written agreement should be kept in good standing as long as LUPIN-TOLVAPTAN treatment is continued.

LUPIN-TOLVAPATN is available for treatment of patients with ADPKD only through a manufacturer and product-specific controlled hepatic safety monitoring and distribution (HSMD) programme conducted and maintained by, or for, the market authorisation holder of LUPIN-TOLVAPATN. A duly signed manufacturer and product-specific PPAF is required for enrollment in the HSMD programme. For more information on the programme, please call. 1-866-488-6017.

Prior to initiation of treatment with LUPIN-TOLVAPTAN, it is important to determine whether expected benefit-risk is deemed to be favourable for the individual patient to be treated. Based on TEMPO 3:4, patients most likely to benefit from tolvaptan appear to be those with rapidly progressive ADPKD, or at a stage of incipient rapid progression, but before widespread destruction of renal architecture has occurred. Factors associated with rapid progression of ADPKD include, large total renal cyst mass for a given age, as measured by total kidney volume (TKV), CKD Stage 2-3, rapid deterioration of renal function, presence of systemic hypertension or albuminuria (also see, [Dosing Considerations](#), below). Conversely, ADPKD patients without evidence of hypertension, and especially those at an early stage of disease with excellent renal function, e.g., estimated creatinine clearance (eCrCL) ≥ 120 mL/min, consistent with renal glomerular (compensatory) hyperfiltration, appear to show little near-term benefit in terms of TKV progression or diminution of renal function decline. From REPRISÉ, patients most likely to benefit from tolvaptan appear to be those at high risk of progressive eGFR decline based on renal function for age (18 to 65 years of age with baseline eGFR between 25 and 65 mL/min/1.73m²).

All patients need to be apprised of the risk of idiosyncratic drug-induced liver injury associated with tolvaptan use in ADPKD (see [WARNINGS AND PRECAUTIONS, Hepatotoxicity](#)), and the need for ongoing monitoring of hepatic function during LUPIN-TOLVAPTAN treatment.

Once LUPIN-TOLVAPTAN treatment is instituted, all patients should be encouraged to drink

water liberally on an ongoing basis in order to match increased urine output, to reduce the likelihood of dehydration and hypernatremia from the aquaretic effects of tolvaptan (see [WARNINGS AND PRECAUTIONS, Dehydration](#)).

Recommended Dose and Dosage Adjustment

LUPIN-TOLVAPTAN is to be administered twice daily in split tolvaptan dose regimens of 45+15 mg, 60+30 mg, or 90+30 mg. According to these split-dose regimens, the total daily tolvaptan doses are 60, 90, or 120 mg, respectively.

The initial dosage for LUPIN-TOLVAPTAN is generally 60 mg tolvaptan per day, as a split-dose regimen of 45+15 mg, with 45 mg taken upon waking, and 15 mg taken approximately 8 hours later. The initial dose should be titrated upward to a split-dose regimen of 90 mg tolvaptan (60+30 mg) per day, and then to a target split-dose regimen of 120 mg tolvaptan (90+30 mg) per day, if tolerated, with at least weekly intervals between titrations. Dose titration should be performed judiciously to ensure that high doses are not poorly tolerated through overly rapid up-titration. Physicians may down-titrate to lower doses based on patient tolerability, and up-titrate again when appropriate. Patients should normally be maintained on the highest tolerated dose of tolvaptan.

The aim of dose titration is to block activity of vasopressin at the renal V_2 receptor as completely and constantly as possible, while maintaining acceptable fluid balance, in order to achieve optimal effects on TKV progression or diminution of renal function decline. The adequacy of vasopressin suppression at a given dose of LUPIN-TOLVAPTAN can be monitored through measurement of urine osmolality and may be used to optimise the clinical benefit of LUPIN-TOLVAPTAN in ADPKD patients. Treatment with tolvaptan is more likely to achieve a better clinical response in patients with greater mean changes from baseline urine osmolality (see [CLINICAL TRIALS, Figures 4 and 5](#)). A target of a decrease of at least 300 mOsm/kg from baseline may be considered ideal in most cases, although a decrease of at least 200 mOsm/kg from baseline may be appropriate in those at moderate risk of disease progression and relatively early in their disease course, but still eligible for tolvaptan treatment. If possible, an absolute urine osmolality of less than 300 mOsm/kg should be maintained at all times (see [CLINICAL TRIALS](#)).

Measurement of urine concentrating ability prior to initiation of LUPIN-TOLVAPTAN will help determine the subsequent level of vasopressin blockade obtained with a particular dosage regimen of tolvaptan. Such subsequent measurements may then be useful in guiding dose titration with tolvaptan, particularly in patients in whom tolerability is dose limiting. Initial assessment of urine concentrating ability before initiation of LUPIN-TOLVAPTAN should be carried out following complete overnight fluid restriction of 10-14 hours, using urine osmolality or specific gravity. During treatment with LUPIN-TOLVAPTAN, measurement of urine osmolality or specific gravity should be carried out at trough before the morning dose, to determine the change in urine osmolality from baseline. However, patients then taking tolvaptan should not be restricted in their usual fluid intake overnight. Although not specifically evaluated, a urine specific gravity in the range of 1.005 generally corresponds to a $U_{osm} < 300$ mOsm/kg, although use of specific gravity measurement is considered less accurate than direct measurement of urine osmolality.

Patients should be told that unnecessary treatment interruption should be avoided, and that daily adherence to the recommended dosing regimen of LUPIN-TOLVAPTAN is important in order to achieve best outcomes in terms of diminution of renal cyst progression and preservation of renal function (see [ACTION AND CLINICAL PHARMACOLOGY, Pharmacodynamics](#)).

LUPIN-TOLVAPTAN may be taken with or without meals.

Therapy should be interrupted if the ability to drink or accessibility to water is limited.

Dosing Considerations

Liver testing must be performed, including total bilirubin, prior to initiating treatment with LUPIN-TOLVAPTAN to establish baseline liver function. LUPIN-TOLVAPTAN is contraindicated in patients with clinically relevant impairment of hepatic function (also, see [WARNINGS AND PRECAUTIONS, Hepatotoxicity](#), and/or [Hepatic Impairment](#)). Treatment with LUPIN-TOLVAPTAN should not normally be initiated in patients with AST/ALT >3 x ULN. However, substantial hepatic cyst burden, without advanced impairment of hepatic function, should not serve as an impediment to initiation of LUPIN-TOLVAPTAN treatment for ADPKD. If, at any time during treatment with tolvaptan, a patient shows abnormal ALT, AST or total bilirubin (BT) levels that fulfil the criteria for permanent discontinuation, tolvaptan treatment should not be reinitiated. In case of abnormal baseline levels below the limits for permanent discontinuation, pre-existing liver disease is likely, and treatment should only be initiated if the potential benefits of treatment outweigh the potential risks and liver function testing must continue at increased frequency. The advice of a hepatologist is recommended or ADPKD specialist.

To mitigate the risk of significant and/or irreversible liver injury, regular and ongoing blood testing for hepatic transaminases is required during LUPIN-TOLVAPTAN treatment, on a monthly basis for the first 18 months, then every 3 months for 12 months, and thereafter at 3-6 month intervals during treatment with LUPIN-TOLVAPTAN.

At the onset of symptoms or signs consistent with hepatic injury, or if abnormal ALT, or AST increases are detected, LUPIN-TOLVAPTAN administration must be immediately interrupted and repeat liver tests, i.e., ALT, AST, total bilirubin, alkaline phosphatase, must be obtained as soon as possible, ideally within 48-72 hours.

Testing should continue at an increased frequency until symptoms/signs/laboratory abnormalities stabilise or resolve, at which point cautious re-initiation of LUPIN-TOLVAPTAN may be considered.

Current clinical practice suggests that LUPIN-TOLVAPTAN therapy should be interrupted upon confirmation of sustained or increasing transaminase levels, and permanently discontinued if significant increases and/or clinical symptoms of hepatic injury persist. Recommended guidelines for permanent discontinuation include:

- ALT or AST >8-times ULN
- ALT or AST >5-times ULN, for more than 2 weeks
- ALT or AST >3-times ULN, **and** total bilirubin > 2 x ULN or INR > 1.5
- ALT or AST >3-times ULN, with persistent symptoms of hepatic injury as noted above.

If ALT and AST levels remain $<3 \times$ ULN, LUPIN-TOLVAPTAN therapy may be cautiously continued with frequent monitoring, as transaminase levels appear to stabilise during continued therapy in some patients without increases in other liver function tests.

Hepatic Impairment

The effect of hepatic impairment on tolvaptan concentrations in treatment of ADPKD has not been studied. These patients should be managed cautiously and liver enzymes should be monitored regularly.

Due to its potential to induce hepatocellular injury, patients with clinically relevant impairment of hepatic function should not be treated with tolvaptan (see [CONTRAINDICATIONS](#), and [WARNINGS AND PRECAUTIONS, Hepatotoxicity](#)).

Renal Impairment

Patients who are anuric, or at, or approaching end-stage-renal disease (ESRD), would not be expected to benefit from tolvaptan treatment (see [WARNINGS AND PRECAUTIONS, Renal Impairment](#), and [ACTION AND CLINICAL PHARMACOLOGY, Renal Insufficiency](#)). Treatment should be avoided in these patients.

Increases in Serum Uric Acid

Tolvaptan use may lead to increases in serum uric acid and gout (see [WARNINGS AND PRECAUTIONS, Hyperuricemia](#), [ADVERSE REACTIONS, Increases in Serum Uric Acid](#), and [ACTION AND CLINICAL PHARMACOLOGY, Pharmacodynamics](#)). Uric acid concentrations should be evaluated prior to initiation of LUPIN-TOLVAPTAN therapy, and as indicated during treatment.

CYP-3A Inhibitors

Concomitant use of LUPIN-TOLVAPTAN with strong CYP-3A inhibitors is contraindicated (see [CONTRAINDICATIONS, DRUG INTERACTIONS](#)).

For patients taking moderate inhibitors of CYP-3A concomitantly (see [WARNINGS AND PRECAUTIONS, CYP-3A Inhibitors](#)), split-dose regimens should be down-adjusted by half, e.g., split dose regimens of 90+30 mg per day should be down-adjusted to 45+15 mg per day; 45 mg/15 mg per day should be adjusted to 15 mg/15 mg per day; 60 mg/30 mg per day should be adjusted to 30 mg/15 mg per day. Further reductions should be considered if patients cannot tolerate the reduced tolvaptan doses. With such down-adjustments, the second daytime dose should be maintained at 15 mg, and the first daily dose down-adjusted, as required.

CYP-3A Inducers

Concomitant use of LUPIN-TOLVAPTAN with strong CYP-3A inducers should be avoided (see [WARNINGS AND PRECAUTIONS, Drugs metabolised by CYP-3A or transported by P-gp](#), and [DRUG INTERACTIONS](#)).

Missed Dose

If a patient misses a dose of LUPIN-TOLVAPTAN, the patient should take their next dose at its scheduled time and prescribed dose level.

Administration

Tolvaptan can be taken without regard to food or the timing of food. It should not be taken with

grapefruit juice, or after eating grapefruit, as this may cause a significant increase in tolvaptan concentrations.

OVERDOSAGE

In healthy subjects, single oral doses of tolvaptan of up to 480 mg, and multiple doses up to 300 mg once daily for 5 days have been well tolerated in trials. There is no specific antidote for tolvaptan intoxication. The signs and symptoms of an acute overdose can be anticipated to be those of excessive pharmacologic effect, that is, a rise in serum sodium concentration, polyuria, thirst, and dehydration/hypovolemia.

No mortality was observed in rats or dogs following single oral doses of 2,000 mg/kg (maximum feasible dose). A single oral dose of 2,000 mg/kg was lethal in mice and symptoms of toxicity in affected mice included decreased locomotor activity, staggering gait, tremor and hypothermia.

If overdose occurs, estimation of the severity of poisoning is an important first step. A thorough history and details of overdose should be obtained, and a physical examination should be performed. The possibility of multiple drug involvement must be considered.

Treatment should involve symptomatic and supportive care, with respiratory, ECG and blood pressure monitoring, and water/electrolyte supplements as needed. A profuse and prolonged aquaresis should be anticipated, which, if not matched by oral fluid ingestion, should be replaced with intravenous hypotonic fluids, while closely monitoring electrolytes and fluid balance.

ECG monitoring should begin immediately and continue until ECG parameters are within normal ranges. Dialysis may not be effective in removing tolvaptan because of its high binding affinity for human plasma protein (> 98%). Close medical supervision and monitoring should continue until the patient recovers.

For the most recent information in the management of a suspected drug overdose, contact your regional poison control centre or Health Canada's toll-free number, 1-844 POISON-X (1-844-764-7669).

ACTION AND CLINICAL PHARMACOLOGY

Mechanism of Action

Tolvaptan is a selective vasopressin V₂-receptor antagonist with an affinity for the V₂-receptor that is 1.8 times that of native arginine vasopressin (AVP). Tolvaptan affinity for the V₂-receptor is 29 times greater than for the V_{1a}-receptor. When taken orally, tolvaptan inhibits the binding of vasopressin at the V₂-receptor in the kidney. The decreased binding of vasopressin to the V₂-receptor lowers adenylate cyclase activity resulting in a decrease in intracellular adenosine 3', 5'-cyclic monophosphate (cAMP) concentrations. In human ADPKD cyst epithelial cells, tolvaptan inhibited AVP-stimulated *in vitro* cyst growth and chloride-dependent fluid secretion into cysts. In animal models, decreased cAMP concentrations were associated with decreases in the rate of growth of total kidney volume and the rate of formation and enlargement of kidney cysts. Inhibition of the V₂-receptor in the kidney epithelial cells also prevents aquaporin 2 containing vesicles from fusing with the plasma membrane, which in turn results in an increase in free water clearance, i.e., aquaresis, and a decrease in urine osmolality.

Tolvaptan metabolites have no or weak antagonist activity for human V₂-receptors, compared with tolvaptan. *In vitro* studies have found both enantiomers of tolvaptan to be equally potent at the V₂ receptor.

Pharmacodynamics

With the recommended split-dose regimens in patients with autosomal dominant polycystic kidney disease (ADPKD), tolvaptan inhibits vasopressin from binding to the V₂-receptor in the kidney for the entire day, as indicated by increased urine output and decreased urine osmolality by about 250-300 mOsm/kg from baseline, or to below 300 mOsm/kg. Higher doses, e.g., 90+30 mg/day, reduced urine osmolality to below 300 mOsm/kg in a greater proportion of patients than the lower doses.

In healthy subjects or patients with CKD Stage 1 to 4 receiving a single dose of tolvaptan, the onset of the aquaretic effects occurs within 1 to 2 hours post-dose. In healthy subjects, single doses of 60 and 90 mg produce a peak effect of about 9 mL/min increase in urine excretion rate that is observed between 4 and 8 hours post-dose. Higher doses of tolvaptan do not increase the peak effect in urine excretion rate but sustain the effect for a longer period of time. The offset of tolvaptan action is rapid with urine excretion rate returning to baseline within 24 hours following a 90 mg dose.

Increases in daily urine output in response to tolvaptan treatment are positively correlated with baseline renal function. Following a 90+30 mg split-dose regimen in patients with CKD Stage 1 or 2, the change in mean daily urine volume was about 4 L for a mean total daily volume of about 7 L. In Stage 4 patients, the mean change in daily urine volume was about 2 L for a total daily urine volume of about 5 L. Urine osmolality appears to be maximally suppressed at a urine excretion rate of about 4 mL/min or about 5 L/day.

The increase in free water excretion can result in an increase in serum sodium concentration unless fluid intake is increased to match urine output. Following a 90+30 mg split-dose regimen of tolvaptan in patients with CKD Stage 1 to 4, mean serum sodium concentrations were increased about 2 mEq/L.

Plasma concentrations of native AVP may increase (avg. 2-9 pg/mL) with tolvaptan treatment and return to baseline levels when treatment is stopped.

With tolvaptan treatment, small decreases in glomerular filtration rate, as measured by iothalamate clearance, in the order of 6-10% are observed soon after tolvaptan initiation, and are independent of baseline renal function. It is noted that percent changes in renal plasma flow are highly correlated to percent changes in GFR, with these changes reversible upon discontinuation of tolvaptan. It is believed these changes may occur in response to the observed decrease in urine osmolality caused by tolvaptan.

Serum concentrations of creatinine and cystatin C are slightly increased in patients with CKD Stage 1 to 3, with creatinine changes 2-fold larger in patients with CKD Stage 4.

Uric acid clearance is decreased about 20% -25% in patients with eGFR_{MDRD}>30 mL/min/1.73m². Mean uric acid values observed at baseline in the pivotal registration trial were 5.6 mg/dL (n= 948), increased to 6.4 mg/dL (n= 907) following 3 weeks of tolvaptan titration, and were 6.5 mg/dL (n= 721) following 36 months of treatment with tolvaptan (see [WARNINGS AND PRECAUTIONS, Hyperuricemia](#)). With placebo, mean uric acid values observed at baseline

were 5.5 mg/dL (n= 482), increased to 5.6 mg/dL (n= 474) following 3 weeks of treatment, and were 5.9 mg/dL (n= 406) following 36 months of treatment.

After 3 weeks of tolvaptan treatment at doses from 90 to 120 mg/day, ADPKD patients demonstrated a total kidney volume (TKV) reduction of approximately 3.7%, $p < 0.0001$ (see [CLINICAL TRIALS, Figure 1](#)). Treatment with tolvaptan appears more likely to achieve a greater response in patients with the greatest mean changes from baseline in urine osmolality (see [CLINICAL TRIALS, Figures 4](#) and [5](#)). Repeated “breakthrough” from inhibition, as indicated by a urine osmolality greater than plasma osmolality, i.e., > 300 mOsm/kg, may provide a stimulus for cyst cell division and progression of the disease.

In healthy subjects, no prolongation of the QT interval was observed with tolvaptan following multiple doses of 300 mg/day for 5 days.

Pharmacokinetics

In healthy subjects, the pharmacokinetics of tolvaptan after single doses of up to 480 mg and multiple doses up to 300 mg once daily have been examined. For single doses, area under the curve (AUC) increases proportionally with dose. After administration of doses ≥ 60 mg, however, C_{max} increases less than proportionally with dose. For multiple administration of 300 mg doses compared to 30 mg doses, C_{max} and AUC were only 4.2- and 6.4-fold higher. The pharmacokinetic properties of tolvaptan are stereospecific, with a steady-state ratio of the S-(-) to the R-(+) enantiomer of about 3.

Following oral administration of tolvaptan, peak concentrations are observed between 2 and 4 hours post- dose.

Moderate or severe hepatic impairment or congestive heart failure decrease the clearance and increase the volume of distribution of tolvaptan.

Absorption:

The absolute bioavailability of tolvaptan is 56% (range 42-80%). *In vitro* data indicate that tolvaptan is a substrate and inhibitor of P-gp.

Ingestion of a single dose of 90 mg of tolvaptan with a high-fat meal increased tolvaptan C_{max} 1.96-fold, with no increase in AUC. Similar intake of 60 mg and 30 mg increased tolvaptan C_{max} 1.4- and 1.2-fold, respectively, with no significant increase in AUC.

Distribution:

Tolvaptan is highly plasma protein bound (98%).

Metabolism:

Tolvaptan is extensively metabolized with less than 1% of the dose excreted unchanged in the urine.

Tolvaptan is a CYP-3A substrate and does not appear to have clinically meaningful inhibitory activity. *In vitro* trials indicated that tolvaptan was extensively metabolized by the cytochrome P450 isoenzyme CYP- 3A4/5 and formed many metabolites, with fourteen identified in plasma, urine and feces. The metabolism of most tolvaptan metabolites was also mediated by CYP- 3A4/5.

Excretion:

Tolvaptan is eliminated entirely by non-renal routes, with 19% of a radioactive dose excreted as unchanged tolvaptan in the feces. The rest is metabolised mainly, if not exclusively, by CYP-3A. After oral dosing, clearance is about 4 mL/min/kg and the terminal phase half-life is about 9 hours. The accumulation factor of tolvaptan with the once-daily regimen is 1.3 and the trough concentrations amount to $\leq 16\%$ of the peak concentrations, suggesting a dominant half-life somewhat shorter than 9 hours. There is marked inter- subject variation in peak and average exposure to tolvaptan with a percent coefficient of variation ranging between 30 and 60%.

Special Populations and Conditions**Pediatrics:**

The pharmacokinetics of tolvaptan in patients under the age of 18 years have not been studied.

Geriatrics:

Age did not substantially influence the pharmacokinetic characteristics of tolvaptan following single-dose or multiple-dose administration of 60 mg tablets.

Gender:

Gender was found to have no significant effect on tolvaptan pharmacokinetics.

Race:

In an open-label crossover trial, 24 Japanese and 25 Caucasian men were administered a single 30 mg oral dose of tolvaptan. The mean tolvaptan C_{max} and AUC_{∞} values were only 5-15% higher in Japanese subjects compared to Caucasian subjects.

Hepatic Insufficiency:

In a population pharmacokinetic analysis of hyponatremia patients, moderate hepatic impairment was associated with a 19% decrease in tolvaptan clearance and severe hepatic impairment was associated with a 24% decrease in clearance and a 50% increase in volume of distribution. The effects of such changes have not been studied in ADPKD patients who typically have normal hepatic function despite having variable degrees of polycystic liver disease.

Renal Insufficiency:

Tolvaptan has been studied in subjects with varying degrees of renal function following a single 60-mg dose. Tolvaptan AUC_{∞} in subjects with $CrCL < 30$ mL/min was approximately 1.9 times higher than that in subjects with $CrCL > 60$ mL/min, but there was no correlation between tolvaptan AUC_{∞} and changes in pharmacodynamic endpoints (urine volume, fluid intake, creatinine and free water clearances, urinary excretions of creatinine, Na^+ and K^+). Tolvaptan increased free water clearance and suppressed urine osmolality to below 300 mOsm/kg in all subjects studied. Increases in urine output were positively correlated with baseline renal function and were significantly lower in patients with CKD Stage 4, i.e., $GFR < 30$ mL/min.

No clinical trial data is available for tolvaptan in patients with a creatinine clearance < 10 mL/min or in patients on chronic dialysis.

STORAGE AND STABILITY

Store LUPIN-TOLVAPTAN at 15 °C to 30 °C.

DOSAGE FORMS, COMPOSITION AND PACKAGING

LUPIN-TOLVAPTAN (tolvaptan) is available in weekly combination blister packs of 15+15 mg, 30+15 mg, 45+15 mg, 60+30 mg, and 90+30 mg tablets.

Each carton contains 4 weekly packs (e.g., 45+15 mg (4 weekly packs each containing 1 blister card with 14 tablets (7 x 45 mg tablets and 7 x 15 mg tablets))).

Each blister pack will contain a total of 14 tablets, 7 tablets of each strength, with one tablet from the row of the higher dose to be taken each morning and one tablet from the row of the lower dose to be taken each evening.

LUPIN-TOLVAPTAN 15 mg tablets are non-scored, Pink to light pink coloured, capsule shape, mottled tablet debossed with “F05” on one side and “LU” on other side

LUPIN-TOLVAPTAN 30 mg tablets are non-scored, Pink to light pink coloured, round shaped, flat faced bevelled edge mottled tablet debossed with “F06” on one side and “LU” on other side.

LUPIN-

TOLVAPTAN 45 mg tablets are non-scored, Pink to light pink coloured, octagonal shape, mottled tablets debossed with “LU” on one side and “F07” on other side

LUPIN-TOLVAPTAN 60 mg tablets are non-scored, Pink to light pink coloured, almond shaped, flat faced bevelled edge mottled tablet debossed with “LU” on one side and “F08” on other side

LUPIN-TOLVAPTAN 90 mg tablets are non-scored, Pink to light pink coloured, capsule shaped, biconvex mottled tablets debossed with “LU” on one side and “F09” on other side.

Inactive Ingredients: Corn starch, dehydrated alcohol, hydroxypropyl cellulose, lactose monohydrate, low substituted hydroxypropyl cellulose, magnesium stearate, microcrystalline cellulose, and Ferric oxide (Red iron), traces of methylene chloride and ethanol maybe present.

PART II: SCIENTIFIC INFORMATION

PHARMACEUTICAL INFORMATION

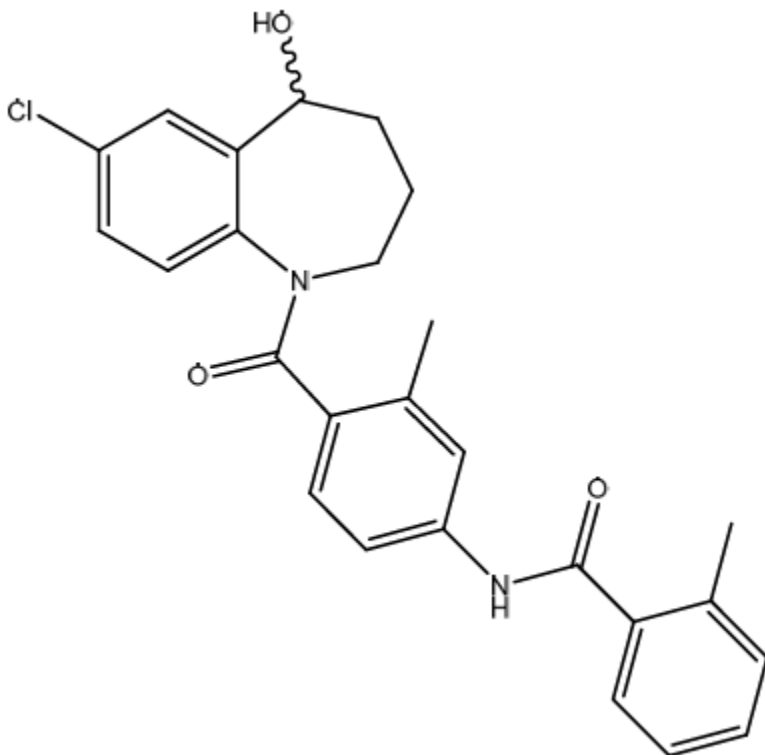
Drug Substance

Common name: Tolvaptan

Chemical name: (\pm)-4'-[(7-chloro-2,3,4,5-tetrahydro-5-hydroxy-1*H*-1-benzazepin-1-yl) carbonyl]-*o*-tolu-*m*-toluidide

Molecular formula and molecular mass: C₂₆H₂₅ClN₂O₃ 448.94

Structural formula:



Physicochemical properties: Tolvaptan is a white crystalline powder. It is practically insoluble in

water (0.00005 w/v% at 25 °C), and no pH dependence of solubility was observed. Tolvaptan is stable to light.

CLINICAL TRIALS

Comparative Bioavailability Studies

A double blind, Balanced, Randomized, Single-Dose, Two-Treatment, Two-Sequence, Two-Period Crossover Oral Bioequivalence Study Comparing Tolvaptan Tablets 90 mg manufactured by Lupin Limited, India, with JINARC (tolvaptan) Tablets 90 mg manufactured by Otsuka Pharmaceutical Co., Ltd., Tokyo, 101-8535 Japan, distributed by Otsuka Canada Pharmaceutical Inc. Saint-Laurent, Quebec H4S 2C9 in Healthy, Adult, Human Male Subjects Under Fasting Conditions.

Comparative bioavailability data from 36 subjects that were included in the statistical analysis are presented in the following table.

SUMMARY TABLE OF THE COMPARATIVE BIOAVAILABILITY DATA

Tolvaptan (1 x 90mg; Fasted study) From measured data Geometric Mean Arithmetic Mean (CV %)				
Parameter	Test¹	Reference²	% Ratio of Geometric Means	Confidence Interval[#]
AUC _T (hr*ng/mL)	7040.6 7943.7 (49.5)	6688.9 7372.6 (46.7)	105.1	97.2 - 113.7
AUC _I (hr*ng/mL)	7242.8 8153.1 (49.3)	6868.0 7548.4 (46.2)	105.3	97.2 - 114.1
C _{MAX} (ng/mL)	625.1 684.8 (43.2)	642.1 675.7 (34.3)	96.7	88.5- 105.7
T _{MAX} ³ (h)	3.1(52.7)	2.7(46.7)	Not applicable	Not applicable
T _{1/2} ³ (h)	8.0 (23.4)	7.5 (19.3)	Not applicable	Not applicable

¹ Lupin-Tolvaptan Tablets 90 mg per tablets

² PrJinarc^{®MD} 90 mg tablets, distributed by Otsuka Canada Pharmaceutical, Canada

³ Expressed as the arithmetic mean (CV%) only

A double blind, Balanced, Randomized, Single-Dose, Two-Treatment, Two-Sequence, Two-Period Crossover Oral Bioequivalence Study Comparing Tolvaptan Tablets 90 mg manufactured by Lupin Limited, India, with JINARC (tolvaptan) Tablets 90 mg manufactured by Otsuka Pharmaceutical Co., Ltd., Tokyo, 101-8535 Japan distributed by Otsuka Canada Pharmaceutical Inc. Saint-Laurent, Quebec H4S 2C9 in Healthy, Adult, Human Male Subjects Under Fed Conditions.

Comparative bioavailability data from 38 subjects that were included in the statistical analysis are presented in the following table.

SUMMARY TABLE OF THE COMPARATIVE BIOAVAILABILITY DATA

Tolvaptan (1 x 90mg; Fed Study) From measured data Geometric Mean Arithmetic Mean (CV %)				
Parameter	Test ¹	Reference ²	% Ratio of Geometric Means	Confidence Interval [#]
AUC _T (hr*ng/mL)	7916.9 8589.8 (43.7)	8345.5 9066.6(46.8)	94.9	88.7 - 101.5
AUC _I (hr*ng/mL)	8044.4 8719.0 (43.6)	8463.9 9194.2(47.0)	95.0	88.8 - 101.7
C _{MAX} (ng/mL)	1004.8 1043.5 (29.2)	1100.4 1149.1 (31.5)	91.3	85.1 - 98.0
T _{MAX} ³ (h)	3.7 (26.8)	3.6 (43.4)	Not applicable	Not applicable
T _{1/2} ³ (h)	5.9 (24.5)	6.0 (29.0)	Not applicable	Not applicable

¹ Lupin-Tolvaptan Tablets 90 mg per tablets

² PrJinarc[®]MD 90 mg tablets, distributed by Otsuka Canada Pharmaceutical, Canada

³ Expressed as the arithmetic mean (CV%) only

The efficacy of tolvaptan in slowing the progression of kidney enlargement (reflecting cyst growth) and kidney function decline is based on randomized, double-blind, placebo-controlled trials in early to late stage of ADPKD.

TEMPO 3:4

In the pivotal double-blind, 36-month, placebo-controlled, multi-center trial, called TEMPO 3:4, a total of 1,445 adult patients (age 18-50 years) with early, rapidly-progressing ADPKD (meeting modified Ravine criteria, total kidney volume (TKV) \geq 750 mL, and estimated creatinine clearance \geq 60 mL/min) were randomised 2:1 to treatment with tolvaptan or placebo, respectively. A total of 1,444 patients were treated for up to 3 years, then followed for 14-42 days after treatment withdrawal. Randomisation was stratified based on several predictors of more rapid progression, namely baseline hypertension status, kidney volume and renal function. All patients remained on standard concomitant medications. Patients were evaluated at screening, baseline, during weekly titration steps and at intervals of at least 4 months for outcome, laboratory and safety assessments. At baseline and yearly visits, magnetic resonance imaging (MRI) of TKV and pharmacokinetic assessments were also performed. Patients who completed the study terminated treatment at 3 years, and were followed for a further period of 2-6 weeks to assess off-drug effects.

Tolvaptan (N=961) and placebo (N=484) groups were well matched, with an average age of 39 years, 52% being male, 84% Caucasian, 13% Asian and 3% other races. At baseline, 79% had hypertension, average estimated glomerular filtration rate (eGFR) was 82 mL/min/1.73 m² calculated by CKD-EPI, and mean TKV was 1,692 mL, with height-adjusted TKV 972 mL/m.

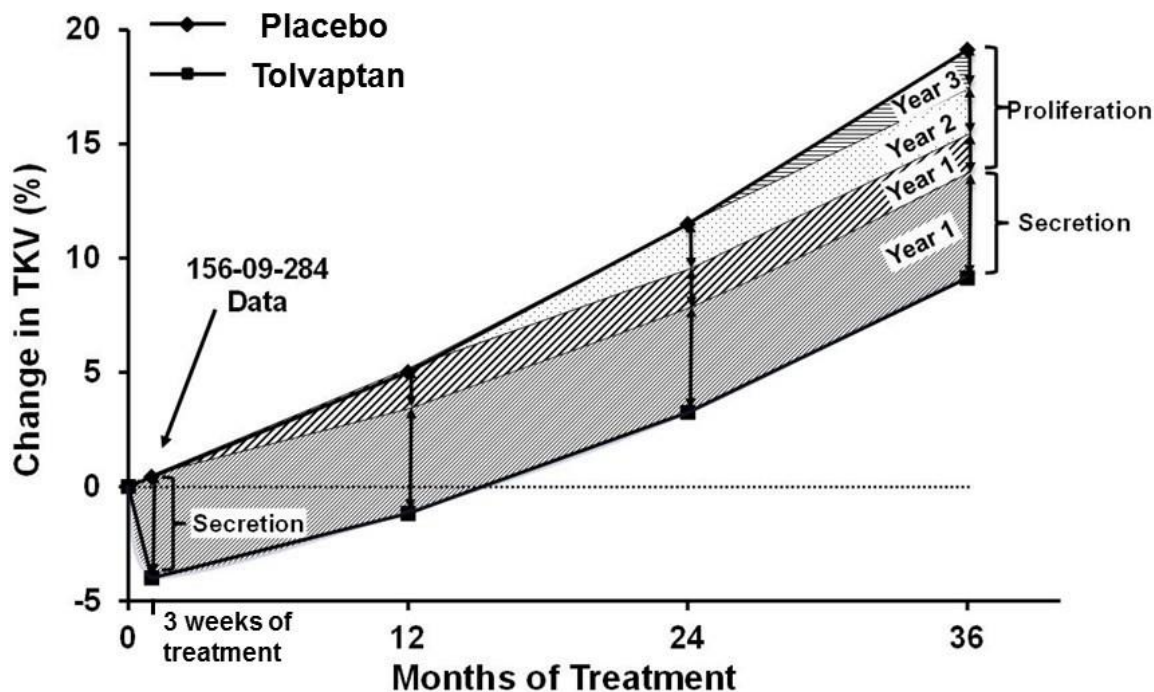
Using $eGFR_{CKD-EPI}$, the distribution of tolvaptan and placebo patients at baseline across CKD stages was as follows: CKD Stage 1 (35%), 2 (48%), and 3 (17%). Within the study population receiving placebo, stratification factors successfully predicted more rapid progression in those who had larger kidneys, lower eGFR, or hypertension at baseline. All patients were encouraged to drink adequate water to avoid thirst or dehydration and at night before retiring.

The primary endpoint was the difference of the rate of change of TKV, normalised as a percentage. TKV increased in the tolvaptan group at a rate of 2.8% per year (95% CI, 2.5% to 3.1%), compared with 5.5% per year (95% CI, 5.1% to 6.0%) for placebo, representing a 49.2% reduction in growth rate averaged over 3 years ($p < 0.0001$).

Analysis also indicated that the effect of treatment on TKV growth was greatest in the first year and included negative cyst growth for the tolvaptan group (-1.7%), compared with positive cyst growth in the placebo group (4.6%), for a treatment effect of -6.3%, a statistically significant difference between groups ($p < 0.0001$). However, tolvaptan has a short-term, so-called “secretory” effect on TKV, presumably due to its aquaretic effect, leading to a diminution of renal cyst fluid, which appears largely reversible upon discontinuation of tolvaptan (see [ACTION AND CLINICAL PHARMACOLOGY, Pharmacodynamics](#)).

This phenomenon will thus apparently overstate the effect of tolvaptan on renal cyst proliferation in the first year when measured by TKV, see Figure 1 below. During the second and third years, kidney enlargement progressed in both groups. Further absolute reductions in TKV growth (relative to placebo) of 1.92% per year (95% CI 2.81 to 1.03%) and 1.78% per year (95% CI 2.77 to 0.78%) were observed at Year 2 and Year 3 of therapy, respectively.

Figure 1: Model of Tolvaptan’s Effects on Total Kidney Volume



This Figure represents analysis of percent change in TKV over time for tolvaptan-treated

patients (intention-to-treat dataset) from TEMPO 3:4, overlaid with 3-week data from a Phase 2 pharmacodynamic study.

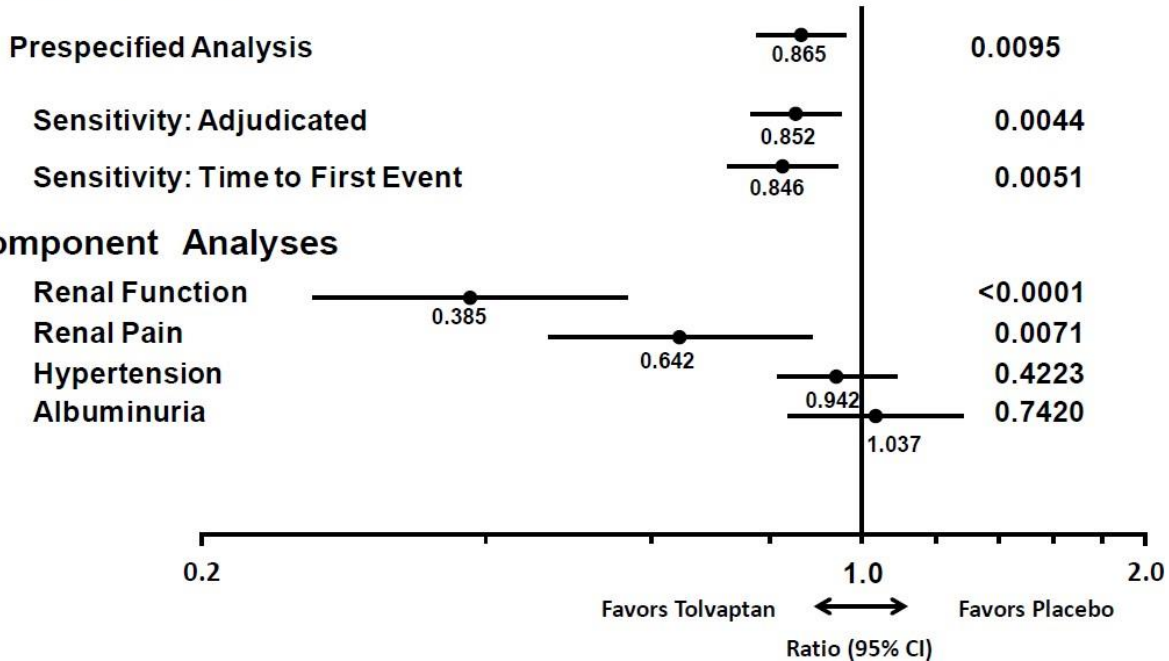
Pre-specified secondary end-points were tested sequentially. The key secondary composite outcome endpoint, reflecting ADPKD progression, was evaluated using a time-to-event analysis, consisting of:

- 1) worsening kidney function, defined as a persistent 25% reduction, i.e., reproduced over at least two weeks, in reciprocal serum creatinine during treatment (equivalent to a 33% increase in serum creatinine), from end of titration to last on-drug visit;
- 2) medically significant kidney pain, defined as requiring prescribed leave, last-resort analgesics, narcotic or anti-nociceptive, radiologic or surgical interventions;
- 3) worsening hypertension, defined as a persistent increase in blood pressure category, or an increase in anti-hypertensive medication(s); or
- 4) worsening albuminuria, defined as a persistent increase in albumin/creatinine ratio category (seen at 2 of 3 successive assessments).

The rate of ADPKD-related events, as assessed by the key secondary composite outcome endpoint, was decreased by 13.5% in tolvaptan-treated patients, compared to placebo-treated patients (44 vs. 50 events; HR 0.87; 95% CI, 0.78 to 0.97; p=0.0095), see Figure 2, below.

Figure 2: Key Secondary Composite Endpoint of Clinical Progression Events in TEMPO 3:4

Composite Analyses



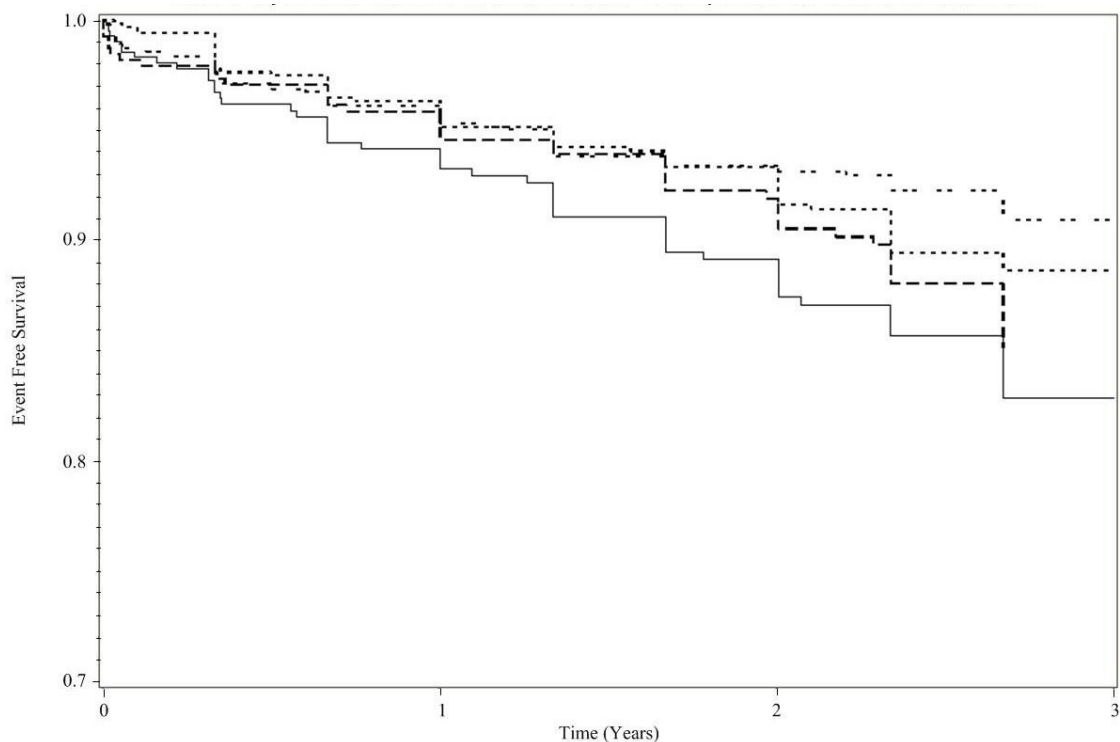
The result of the key secondary composite outcome endpoint is primarily attributed to effects on worsening kidney function and medically significant kidney pain. The renal function events were seen to occur 61.4% less often for tolvaptan compared with placebo (HR 0.39; 95% CI, 0.26 to 0.57; nominal p <0.0001), and renal pain events occurred 35.8% less often in tolvaptan-treated

patients (HR 0.64; 95% CI, 0.47 to 0.89; nominal p=0.007), see Figure 2, above. In contrast, there was no effect of tolvaptan on either progression of hypertension or albuminuria. It should be noted, however, that 79% of patients were already hypertensive at baseline in both groups.

The next sequentially ordered secondary endpoint was the slope of kidney function decline, assessed as change in estimated glomerular filtration rate (eGFR_{CKD-EPI}) during treatment from end of titration to last on-drug visit. The tolvaptan-treated patients had a 26.4% reduction in the rate of renal function decline, compared to placebo, -2.7 versus -3.6 mL/min/1.73m²/year, respectively, p <0.0001. Note that initiation of tolvaptan is associated with a generally reversible, prompt decline in GFR, likely due to hemodynamic factors (see [ACTION AND CLINICAL PHARMACOLOGY, Pharmacodynamics](#)).

Preliminary analyses for urine osmolality (Uosm) suggest that the greatest degree of efficacy, as related to outcomes of events of worsening renal function or renal pain, is seen when Uosm was suppressed by at least -300 mOsm/kg. A lesser degree of efficacy (when compared with other groups of Uosm suppression) is seen for the group between -105 and -300 mOsm/kg, see Figure 3 below.

Figure 3: Time to Events of Worsening Renal Function or Kidney Pain, derived from the Key Secondary Composite Endpoint, in TEMPO 3:4

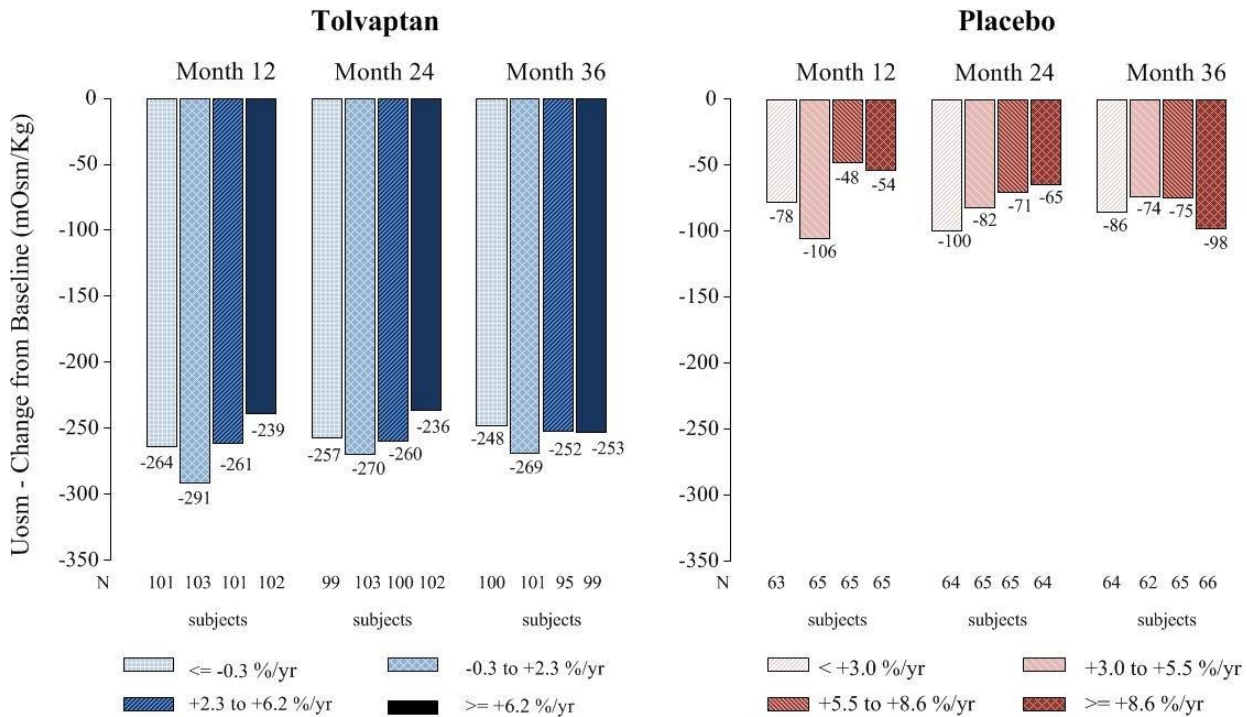


Change of urine osmolality from End of Titration/Week 3 to Baseline; - - - - (-1119 to -300 mOsm/kg),(-300 to -105 mOsm/kg), - - - (-105 to 0 mOsm/kg), and — (0 to +545 mOsm/kg)

Patients treated with tolvaptan demonstrating greater changes in urine osmolality from baseline appeared to have better responses in terms of slowing progression of TKV and renal function decline, see Figures 4 and 5, respectively, below. Note that best responses are represented by the

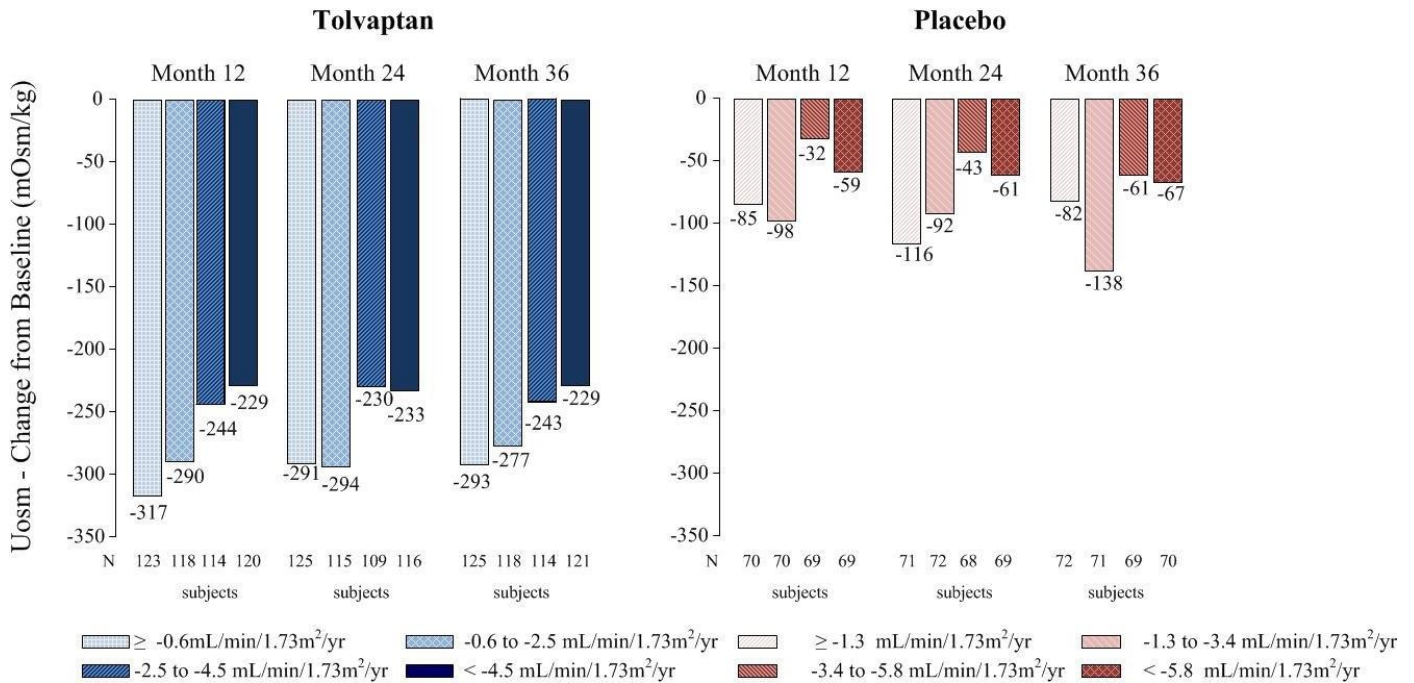
lightest coloured bars to the left, and worst responses in darkest coloured bars presented to the right, for both endpoints of TKV and renal function decline in all tolvaptan and placebo panels. Although a threshold effect seems to have been reached for TKV with any tolvaptan treatment, with all quartiles achieving a mean change in urine osmolality from baseline of at least 200 mOsm/kg and no apparent correlation of relative changes of urine osmolality on TKV growth, there is an apparent correlation of change of urine osmolality from baseline and attenuation of renal function decline, see Figure 5, below. Placebo shows no such effect.

Figure 4: Mean Change from Baseline in Urine Osmolality in TEMPO 3:4, by Quartile of TKV Growth Rate



Quartiles of TKV Growth Rates, ranging from < -0.3 %/year growth to ≥ 6.2 %/year growth for tolvaptan, and from < 3.0 %/year growth to ≥ 8.6 %/year for placebo.

Figure 5: Mean Change from Baseline in Urine Osmolality in TEMPO 3:4, by Quartile of Renal Function Decline, as Measured by Slope of eGFRCKD-EPI



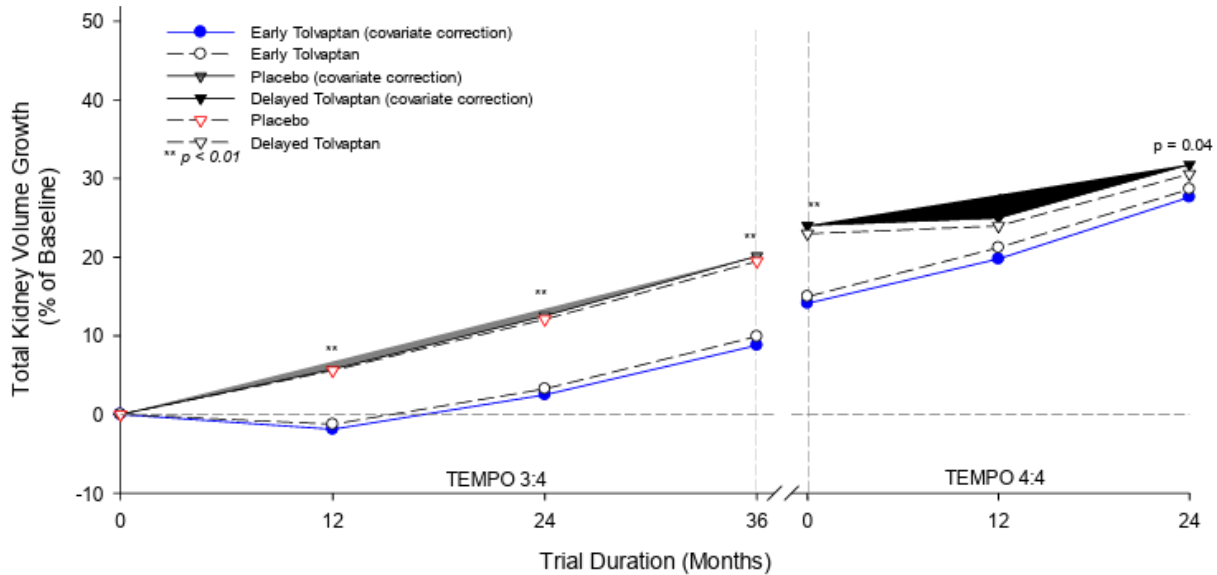
Quartiles of Renal Function Decline, ranging from < 0.6 mL/min/1.73m²/year to > 4.5 mL/min/1.73m²/year of renal function decline for tolvaptan, and from < 1.3 mL/min/1.73m²/year to > 5.8 mL/min/1.73m²/year of renal function decline for placebo

TEMPO 4:4

Following completion of TEMPO 3:4, and after a washout period, most placebo-treated and tolvaptan-treated patients were offered ongoing tolvaptan treatment in TEMPO 4:4, an open-label extension study.

This trial evaluated the effects of tolvaptan on safety, TKV and eGFR in subjects receiving active treatment for 5 years (early-treated), compared with subjects treated with placebo for 3 years, then switched to active treatment for 2 years (delayed-treated). Analysis of TKV progression and eGFR decline for the completed trial are presented below in Figures 6 and 7, respectively.

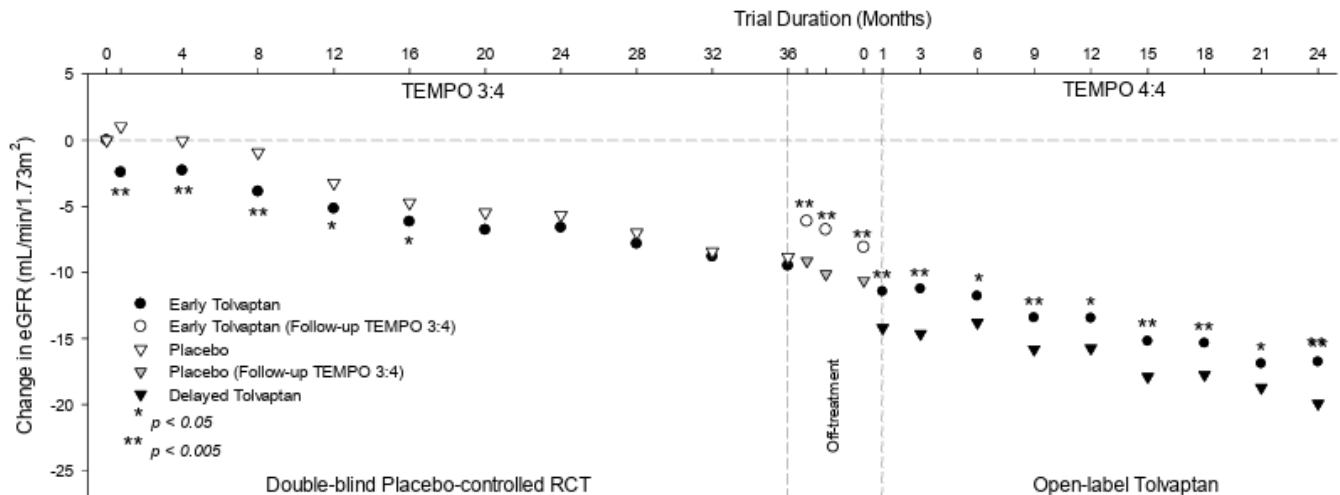
Figure 6: Percent Change in Total Kidney Volume (TKV) from Baseline, from TEMPO 3:4 AND TEMPO 4:4



Average TKV change from baseline for patients receiving tolvaptan (solid circles) or placebo (open triangles) during the three years of placebo-controlled TEMPO 3:4 are shown, followed by a washout period post-treatment (open circles or triangles), and then, for the previously tolvaptan-treated patients, two further years of treatment with tolvaptan in the open-label extension study, TEMPO 4:4 (solid circles for early-treated patients, solid triangles for delayed-treated patients).

The primary endpoint for TKV did not distinguish a difference in change (-1.7%) over the 5-year treatment between early- and delayed-treated subjects at the pre-specified threshold of statistical significance (p=0.3580). Both groups' TKV growth trajectory was slowed, relative to placebo in the first 3 years, suggesting both early- and delayed- tolvaptan treated subjects benefitted to a similar degree.

Figure 7: Change in Renal Function from Baseline, from TEMPO 3:4 (at the left of the Off-Treatment period) and TEMPO 4:4 (at the right of the Off-Treatment period)



Average eGFR change from baseline for patients receiving tolvaptan (solid circles) or placebo (open triangles)

during the three years of placebo-controlled TEMPO 3:4 are shown, followed by off-treatment measurements for approximately 3 months post-treatment (open circles or triangles), and then, for the patients previously treated with tolvaptan in TEMPO 3:4, two further years of treatment with tolvaptan in the open-label extension study, TEMPO 4:4 (solid circles for early-treated patients, solid triangles for delayed-treated patients).

The early-treated subjects experienced a greater overall benefit in renal function than the delayed-treated subjects based on eGFR change over the 5-year treatment period ($p=0.0003$).

A secondary endpoint testing the persistence of positive effects on renal function indicated that the preservation of eGFR observed by the end of the TEMPO 3:4 pivotal trial (3.01 to 3.34 mL/min/1.73m² at follow-up visits 1 and 2) could be preserved during open-label treatment. This difference was maintained in the pre-specified MMRM analysis (3.15 mL/min/1.73m², 95% CI 1.462 to 4.836, $p=0.0003$) and with sensitivity analyses where baseline eGFR data were carried forward (2.64 mL/min/1.73m², 95% CI 0.672 to 4.603, $p=0.0086$). This trial yielded data which supported the long-term effects of tolvaptan on both renal function and TKV, including a demonstrable, incremental renal function benefit for those receiving treatment earlier. These data suggest that tolvaptan can slow the rate of renal function decline, and that these benefits persist over the duration of therapy.

Following an additional 2 years of tolvaptan treatment, resulting in a total of 5 years on tolvaptan therapy, the safety and tolerability were supportive of the results observed in the double-blind, placebo-controlled study.

REPRISE

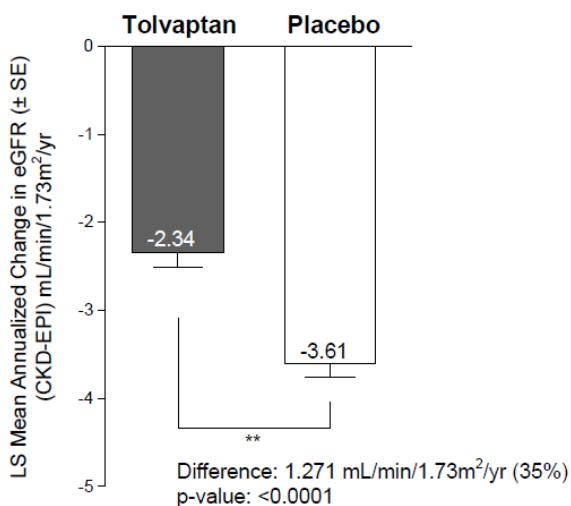
The pivotal double-blind, placebo-controlled, multi-center trial, called REPRISE, included a total of 1,519 adult patients (age 18-65) with chronic kidney disease (CKD) from late stage 2 to early stage 4 (eGFR between 25 and 65 mL/min/1.73m² if younger than age 56; or eGFR between 25 and 44 mL/min/1.73m², plus eGFR decline >2.0 mL/min/1.73m²/year if between age 56-65). They entered an 8-week single-blind pre-randomization phases (screening, placebo run-in, tolvaptan titration and tolvaptan run-in). Tolvaptan was initiated during the single-blind pre-randomization phase and up-titrated every 3-4 days from a daily oral dose of 30 mg/15 mg to 45 mg/15 mg, 60 mg/30 mg and up to a maximum dose of 90 mg/30 mg. Only patients who could tolerate the two highest doses of tolvaptan (60 mg/30 mg or 90 mg/30 mg) were randomized 1:1 to treatment with tolvaptan or placebo. A total of 1,370 patients were randomized and treated during the 12-month double-blind period. After completion of treatment the patients entered a 3-week follow-up period.

Randomization was stratified by patients' baseline characteristics of eGFR, age, and TKV (if available) and all patients remained on standard concomitant medications. Patients were maintained on their highest tolerated dose for a period of 12 months but could interrupt, decrease and/or increase as clinical circumstances warranted within the range of titrated doses. All patients were encouraged to start drinking an adequate amount of water at screening and continuing through the end of the trial to avoid thirst or dehydration. Patients were evaluated weekly during the pre-randomization phases and monthly during randomized treatment for efficacy and safety assessments.

Patients were assigned to tolvaptan (N=683) or placebo (N=687) and were distributed across CKD stages as follows: Stage 2 (5.2%), Stage 3a (30%), Stage 3b (45.1%) and Stage 4 (19.5%). The average age was 47 years.

The primary endpoint (eGFR change) was the treatment difference in the change of eGFR from pre-treatment baseline to post-treatment follow-up, annualized by dividing by each subject's treatment duration. Decline in renal function was 35% slower with tolvaptan compared to placebo ($p < 0.0001$); the change of eGFR from pretreatment baseline to post-treatment follow-up was $-2.34 \text{ mL/min/1.73 m}^2/\text{year}$ with tolvaptan vs $-3.61 \text{ mL/min/1.73 m}^2/\text{year}$ with placebo, with a treatment effect of $1.271 \text{ mL/min/1.73 m}^2/\text{year}$, see as illustrated in Figure 8. The key secondary endpoint (eGFR slope) and all sensitivity analyses were significant (at $p \leq 0.0005$).

Figure 8: Annualized Change in eGFR (CKD-EPI) from Pre-treatment Baseline to Post-treatment Follow-up in REPRISE



Subgroup analysis for the primary and key secondary endpoints demonstrated consistent treatment effects of tolvaptan for subjects in CKD Stages 2, 3a, 3b, and 4.

The results of REPRISE, therefore, supplement the earlier finding that tolvaptan reduces the progression of the enlargement of kidney volume and show that tolvaptan also reduces the rate of regression of renal function.

DETAILED PHARMACOLOGY

Arginine vasopressin (AVP) is a neuropeptide hormone which causes vasoconstriction via V_{1a} -receptors and promotes water reabsorption in the kidneys via V_2 -receptors, both of which are G-protein-coupled transmembrane receptors. The V_2 -receptors are primarily responsible for the anti-diuretic effects of AVP. Patients with various disorders, including congestive heart failure, liver cirrhosis, and syndrome of inappropriate secretion of anti-diuretic hormone (SIADH), are at risk of experiencing excess water retention or inadequate water disposal due to increased vasopressin secretion. Tolvaptan is a competitive, non-peptide vasopressin antagonist drug that blocks the binding of arginine vasopressin at the V_2 -receptors of the distal nephron, thereby inducing water diuresis (aquaresis), but notably without the depletion of electrolytes.

Pharmacodynamics

In vitro antagonistic effects of tolvaptan were investigated in binding experiments using a human endocervical carcinoma cell line (HeLa cells) expressing human AVP receptor subtypes (V_{1a} , V_{1b} , and V_2). Tolvaptan inhibited [^3H]AVP binding to the V_2 -receptors in a concentration-

dependent manner, with an inhibition constant (K_i) of 0.43 ± 0.06 nM, which was approximately 1.8 times higher than that of AVP ($K_i = 0.78 \pm 0.08$ nM). Tolvaptan also inhibited [3 H]AVP binding to the V_{1a} -receptors with a K_i of 12.3 ± 0.8 nM, but the affinity is approximately 29 times weaker than that for V_2 -receptors. On the other hand, tolvaptan did not inhibit [3 H]AVP binding to the V_{1b} -receptors even at 100 nM.

The affinities of tolvaptan for rat and canine AVP receptors were investigated by measuring the inhibition of [3 H]AVP binding to membrane preparations prepared from rat liver (V_{1a}), canine platelet (V_{1a}), and rat and canine kidney (V_2). Tolvaptan concentration-dependently inhibited [3 H]AVP binding to rat AVP V_{1a} - and V_2 -receptors with K_i of 345 ± 54 nM and 1.33 ± 0.26 nM, and to canine AVP V_{1a} - and V_2 -receptors with K_i of 40.3 ± 12.0 nM and 0.66 ± 0.09 nM, respectively. Thus, tolvaptan was approximately 259- and 61-times more selective for V_2 -receptors than for V_{1a} -receptors in rats and dogs.

TOXICOLOGY

Single-Dose Toxicity

Single-dose toxicity trials of tolvaptan were conducted by the oral route at doses of up to 2,000 mg/kg in Sprague-Dawley rats and Beagle dogs. No mortality or clinical signs indicative of toxicity were observed in rats or dogs. The minimum lethal dose was not determined in the single-dose trials in rats or dogs. There were no apparent gender differences in sensitivity to the acute effects of tolvaptan in either rats or dogs.

There was no macroscopic evidence of target organ toxicity at any dose level.

Repeat-Dose Toxicity

Repeat-dose oral toxicity trials were conducted in Sprague-Dawley rats and Beagle dogs for up to 26 weeks and 52 weeks, respectively. In rats, the No Observed Adverse Effect Level (NOAEL) was 1,000 mg/kg/day in both sexes in the 4-week and 13-week repeated oral dose toxicity trials. In the 26-week trial at doses of 30, 100 and 1,000 mg/kg/day, the results showed neither overt toxicity nor target organ toxicity even at 1,000 mg/kg/day. However, 3 females given 1,000 mg/kg/day were euthanized in a moribund state (dehydration). Therefore, the NOAEL in this trial was estimated to be 1,000 mg/kg/day in the males and 100 mg/kg/day in the females (serum drug concentration at the fourth week of administration: C_{max} was 1.37 and 3.42 mcg/mL and AUC_{0-24h} was 12.72 and 20.76 mcg.h/mL in the males and females, respectively). In dogs, the NOAEL was 1,000 mg/kg/day in both sexes in the 4-week and 13-week administration. In the 52-week administration trial at doses of 30, 100 and 1,000 mg/kg/day, the results showed no notable target organ toxicity even at 1000 mg/kg/day. However, one male and 2 females given 1,000 mg/kg/day were sacrificed in a moribund state because of decreases in body weight and food consumption. Therefore, the NOAEL in this trial was estimated to be 100 mg/kg/day in both sexes (serum drug concentration at 52nd week of administration: C_{max} was 5.46 and 6.05 mcg/mL and AUC_{0-24h} was 31.45 and 42.35 mcg.h/mL in the males and females, respectively).

Genotoxicity

The genotoxic potential of tolvaptan was assessed in a battery of *in vitro* and *in vivo* test systems. Tolvaptan exhibited no genotoxic potential at concentrations up to 5,000 mcg/plate in the bacterial (*Salmonella typhimurium* and *Escherichia coli*) reverse-mutation test, up to 200 mcg/mL in the forward gene mutation test in mouse lymphoma cells, following pulse treatment

at up to 100 mcg/mL (in the presence or absence of metabolic activation) or with continuous treatment at up to 40 mcg/mL (in the presence or absence of metabolic activation) in the chromosomal aberration test using Chinese hamster lung fibroblast cell line (CHL) or at doses up to 2,000 mg/kg in rat micronucleus test using bone marrow cells from male and female rats administered tolvaptan.

Carcinogenicity

The carcinogenic potential of tolvaptan was evaluated in one 104-week oral carcinogenicity trial in mice and one 104-week oral carcinogenicity trial in rats. Dose levels for the mouse carcinogenicity trial were 0, 10, 30 and 60 mg/kg/day in males and 0, 10, 30 and 100 mg/kg/day in females. The dose levels for the rat carcinogenicity trial were 0, 100, 300 and 1000 mg/kg/day in males and 0, 30, 100, 300 and 1,000 mg/kg/day in females. Oral (gavage) administration of tolvaptan to B6C3F1 mice or Sprague-Dawley rats for 104 weeks was not associated with a decrease in survival or an increase in the incidence of neoplastic or non-neoplastic drug-related findings in males or females. The highest doses tested in mice resulted in exposures (AUC_{0-24h}) that were just below (females; 4.3317 mcg.h/mL) and below (males; 2.8595 mcg.h/mL) the exposure in humans at the MRHD. (120 mg). The highest dose tested in rats resulted in exposures (AUC_{0-24h}) that were approximately 2-times (males; 12.716 mcg.h/mL) and 5-times (females; 33.449 mcg.h/mL) greater than the exposure in humans at steady state at the MRHD of 120 mg.

Reproductive and Developmental Toxicity Trials

In trials of effects on fertility and reproductive performance in Sprague-Dawley rats, tolvaptan did not impair reproductive performance at doses up to 1,000 mg/kg/day in males and 100 mg/kg/day in females (approximately 81- and 8-times the MRHD on a mg/m² basis, respectively). Fertility was not affected at 1,000 mg/kg/day in males and females. The drug-related effect of altered estrous cycles due to prolongation of diestrus was observed in females given 300 and 1,000 mg/kg/day. The NOAEL was less than 100 mg/kg/day for general toxicologic effects in males and females, 100 mg/kg/day for reproductive performance in females and 1,000 mg/kg/day for reproductive performance in males and for fetal development.

In trials of embryo-fetal development, tolvaptan did not cause any developmental toxicity in rats at maternal doses up to 100 mg/kg/day (8-times the MRHD on a mg/m² basis) or in New Zealand White rabbits at maternal doses up to 300 mg/kg/day (49-times the MRHD on a mg/m² basis). Dose-dependent maternal toxicity was evident in rats at 100 mg/kg/day and higher and in rabbits at 30 mg/kg/day and higher. The NOAEL in rats was 10 mg/kg/day for general toxicologic effects in the parental generation (F₀) dams, 1,000 mg/kg/day for reproductive performance in F₀ dams and 100 mg/kg/day for embryo-fetal development in first generation (F₁) fetuses. Maternal toxicity in female rats consisted of decreased food consumption and body weight (100 mg/kg/day and higher) and developmental toxicity of the F₁ fetuses consisted of decreased body weight and delayed ossification (1,000 mg/kg/day).

In rabbits, the NOAEL was 10 mg/kg/day for general toxicologic effects in F₀ dams, and 100 mg/kg/day for reproductive performance in F₀ dams and 300 mg/kg/day for embryo-fetal development in F₁ fetuses.

Maternal toxicity in female rabbits consisted of decreased food consumption and body weight

(30 mg/kg and higher). In addition, maternal changes in physiology were examined in dams given 1,000 mg/kg and the changes included increased urine volume, decreased urine osmolality, increased water consumption and increased plasma sodium and chloride concentration as well as plasma osmolality and plasma AVP levels. Maternal reproductive performance, as assessed by the ability to maintain pregnancy, was altered at dose levels of 300 mg/kg and higher where a dose-dependent incidence of abortion was observed. There was also evidence of developmental toxicity in rabbits at the maternally toxic dose of 1,000 mg/kg (162-times the MRHD on a mg/m² basis). This developmental toxicity consisted of increased incidences of embryo- fetal death, microphthalmia, open eyelids, cleft palate, brachymelia (zygopodium malformations) and fused phalanx.

Teratogenicity of tolvaptan was further investigated in rabbits. The sensitive period of teratogenicity was during gestation Days 6 to 11, and the maximum sensitivity was shown to be during gestation Days 9 to 11. A toxicokinetics trial in pregnant rabbits showed that 13-day repetitive administration of tolvaptan caused a decrease in exposure level (AUC) of unchanged compound to approximately 1/10 of that at the first administration.

In the prenatal and postnatal trial in pregnant rats, tolvaptan had no effect on offspring development at doses up to 100 mg/kg/day following oral administration to pregnant rats from gestation Day 7 through lactation Day 21. Increased perinatal death and decreased body weight of F₁ generation animals during the lactation period and after weaning were observed in the 1,000 mg/kg/day group. The NOAEL was less than 10 mg/kg/day for general toxicologic effects in F₀ dams, and 1,000 mg/kg/day for reproductive performance in F₀ dams and 100 mg/kg/day in terms of effects on offspring development.

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PART III: CONSUMER INFORMATION**LUPIN-TOLVAPTAN****tolvaptan tablets**

This leaflet is part III of a three-part "Product Monograph" published when LUPIN-TOLVAPTAN was approved for sale in Canada and is designed specifically for Consumers. This leaflet is a summary and will not tell you everything about LUPIN-TOLVAPTAN. Contact your doctor, nurse or pharmacist if you have any questions about the drug.

ABOUT THIS MEDICATION**What the medication is used for:**

LUPIN-TOLVAPTAN is used in adults to treat autosomal dominant polycystic kidney disease (ADPKD).

ADPKD is a genetic disorder which leads to the growth of multiple cysts (sacs filled with fluid) on the kidneys. Over time, the cysts grow in size, which can damage the way your kidneys work, and may ultimately lead to kidney failure.

LUPIN-TOLVAPTAN should only be prescribed by a doctor experienced in the diagnosis and treatment of ADPKD.

What it does:

LUPIN-TOLVAPTAN works by blocking the effects of a hormone called vasopressin, which promotes cyst growth in the kidneys in patients with ADPKD. By blocking the effects of the hormone, LUPIN-TOLVAPTAN is able to reduce the growth of the cysts in the kidneys and slow the decline of kidney function. This should help protect your kidneys from damage and failure.

Because of the way that LUPIN-TOLVAPTAN works, you will produce more urine causing you to urinate more frequently during the day and at night. This may become less pronounced over time.

Before starting this medication:

Before taking LUPIN-TOLVAPTAN you should discuss carefully with your doctor if this medication is suitable for you, taking into consideration the potential benefits and risks involved. Once you and your doctor have agreed that LUPIN-TOLVAPTAN is suitable for you, your doctor will ask that you sign a patient-prescriber agreement form (PPAF), stating that you understand the benefits and risks of treatment and that you agree to take blood tests as prescribed by your doctor to start and remain on treatment. For more information on LUPIN-TOLVAPTAN controlled hepatic safety monitoring and distribution programme, please call 1-866-488-6017.

When it should not be used:

LUPIN-TOLVAPTAN Product Monograph

Do not take LUPIN-TOLVAPTAN if:

- you have been asked to permanently discontinue tolvaptan in the past
- you are allergic to tolvaptan, benzazepine or benzazepine derivatives (e.g. REMERON®), or any non-medicinal ingredients in the formulation.
- you cannot replace fluids by drinking or you can not feel if you are thirsty
- you have elevated sodium (salt) in your blood
- you have a condition associated with a low blood volume
- your liver is not functioning properly and in a way that is of concern to your doctor. The presence of hepatic cysts in and of themselves should not be an impediment to initiation of LUPIN-TOLVAPTAN treatment.
- you are pregnant or plan to become pregnant. It is not known if LUPIN-TOLVAPTAN will harm your unborn baby.
- you are breast-feeding. It is not known if tolvaptan passes into your breast milk. You and your healthcare provider should decide if you will take LUPIN-TOLVAPTAN or breast-feed. You should not do both.
- your body is not able to make any urine. LUPIN-TOLVAPTAN will not help your condition.
- you take certain medicines. These medicines could cause you to have too much LUPIN-TOLVAPTAN in your blood:
 - the antibiotic medicines, clarithromycin or telithromycin
 - the antifungal medicines, ketoconazole or itraconazole
 - the anti-HIV medicines, ritonavir, indinavir, nelfinavir, and saquinavir
 - the antidepressant medicine, nefazodone
- you have one of the following rare hereditary diseases:
 - Galactose intolerance
 - Lapp lactase deficiency
 - Glucose-galactose malabsorption because lactose is a non-medicinal ingredient in LUPIN-TOLVAPTAN.

What the medicinal ingredient is:

tolvaptan

What the non-medicinal ingredients are:

Corn starch, dehydrated alcohol, hydroxypropyl cellulose, lactose monohydrate, low substituted hydroxypropyl cellulose, magnesium stearate, microcrystalline cellulose, and ferric oxide (Red iron), traces of methylene chloride and ethanol maybe present.

What dosage forms it comes in:

Weekly combination blister packs of 15+15 mg, 30+15 mg, 45+15 mg, 60+30 mg, and 90+30 mg tablets.

WARNINGS AND PRECAUTIONS

Serious Warnings and Precautions – Liver damage

Your doctor will arrange blood tests before you start treatment with LUPIN-TOLVAPTAN and then at regular intervals during treatment to check for changes in your liver function.

To prevent pregnancy while on LUPIN-TOLVAPTAN treatment, women of childbearing potential must have effective contraceptive measures in place before and during treatment.

BEFORE you use LUPIN-TOLVAPTAN talk to your doctor, nurse or pharmacist if you:

- have difficulty urinating or have an enlarged prostate
- are dehydrated or suffer from excessive vomiting, diarrhea or sweating
- have low levels of sodium in your blood
- have high potassium levels in your blood
- have high levels of uric acid in your blood or gout
- are taking medication to treat high blood pressure
- are less than 18 years old
- have been asked to permanently discontinue tolvaptan in the past

Driving and using machines: Before doing tasks which require special attention, wait until you know how you respond to LUPIN-TOLVAPTAN. Dizziness, weakness and fainting can occur.

INTERACTIONS WITH THIS MEDICATION

As with most medicines, interactions with other drugs are possible. Tell your doctor, nurse, or pharmacist about all the medicines you take, including drugs prescribed by other doctors, vitamins, minerals, natural supplements, or alternative medicines.

The following may interact with LUPIN-TOLVAPTAN: clarithromycin, ketoconazole, ritonavir, saquinavir, rifampicin, phenytoin, carbamazepine, St. John’s Wort, cyclosporin, quinidine, verapamil, erythromycin, fluconazole, hypertonic saline solutions, drugs that increase serum sodium concentrations, drugs known to increase serum potassium levels (such as spironolactone), digoxin, consuming grapefruit juice, or vasopressin drugs, such as desmopressin, used to control bleeding.

PROPER USE OF THIS MEDICATION

Always take LUPIN-TOLVAPTAN exactly as your doctor has prescribed. Check with your doctor, nurse or *LUPIN-TOLVAPTAN Product Monograph*

pharmacist if you are not sure.

You can take LUPIN-TOLVAPTAN with or without food.

While taking LUPIN-TOLVAPTAN you will produce more urine, urinate more frequently, and urinate at night. If this causes you concern, consult your doctor, nurse or pharmacist. This side-effect should become less pronounced with time.

To prevent becoming dehydrated, have water available to drink at all times while taking LUPIN-TOLVAPTAN. Unless your doctor tells you otherwise, drink plenty of water during the day and one or two glasses before going to bed. If you suffer from vomiting, diarrhea or any other condition that might cause you to become dehydrated while taking LUPIN-TOLVAPTAN call your doctor.

Blood tests:

During treatment with LUPIN-TOLVAPTAN, your doctor will arrange blood tests to check for changes in your liver function, as follows:

- before starting treatment with LUPIN-TOLVAPTAN
- every month for the first 18 months, then
- every 3 months thereafter during treatment for the next 12 months, and every 3-6 months thereafter during treatment

Usual adult dose:

- LUPIN-TOLVAPTAN is to be taken twice a day in two different doses. There are three possible dose combinations that your doctor may prescribe:
 - 45+15 mg, for a total daily dose of 60 mg, or
 - 60+30 mg, for a total daily dose of 90 mg, or
 - 90+30 mg, for a total daily dose of 120 mg.
- Take one tablet of the higher dose (45 mg, 60 mg or 90 mg) in the morning.
- 8 hours later take one tablet of the lower dose (15 mg or 30 mg).
- This schedule was designed to give you the best balance between the amount of medicine in your body and possible side effects (especially night time urination).
- Do not drink grapefruit juice during treatment with LUPIN-TOLVAPTAN. This could cause you to have too great of a drug effect, while taking LUPIN-TOLVAPTAN.

Adjusted dose if you are taking certain medicines:

Your doctor will tell you to reduce your usual dose of LUPIN-TOLVAPTAN if you take certain medicines as these medicines could cause you to have too much LUPIN-TOLVAPTAN in your blood. Your doctor may reduce your dose to the following:

- 15 mg + 15 mg, for a total daily dose of 30 mg.
or
- 30 mg +15 mg, for a total daily dose of 45 mg.

Overdose:

If you think you, or a person you are caring for, have taken too much LUPIN-TOLVAPTAN, contact a healthcare professional, hospital emergency department, regional poison control centre or Health Canada's toll-free number, 1-844 POISON-X (1-844-764-7669) immediately, even if there are no signs or symptoms.

Missed Dose:

Do not miss or skip doses of LUPIN-TOLVAPTAN. If you miss a dose, take your next dose at the scheduled time and prescribed level. Do not take 2 doses at the same time.

SIDE EFFECTS AND WHAT TO DO ABOUT THEM

Side effects may include:

- Thirst
- Increased amount of urine
- Increased frequency of urination during the day and at night
- Headache
- Constipation, diarrhea, dry mouth, indigestion, decreased appetite
- Fatigue, weakness, dizziness
- Trouble sleeping
- Muscle spasms
- Rash, itching
- Abdominal pain

If any of these affects you severely, tell your doctor, nurse or pharmacist.

Serious side effects and what to do about them				
Symptom / effect		Talk to your healthcare professional		Stop taking drug and get immediate medical help
		Only if severe	In all cases	
Common	Increased levels of potassium in the blood: irregular heart beats, muscle weakness and generally feeling unwell		√	

Serious side effects and what to do about them				
Symptom / effect		Talk to your healthcare professional		Stop taking drug and get immediate medical help
		Only if severe	In all cases	
	Increased levels of uric acid in the blood/Gout: joint pain, commonly in the big toe, followed by redness, swelling, warmth		√	
Uncommon	Dehydration: increased thirst, dry mouth and/or skin, fatigue, decreased amount of urine, headache, dizziness, irregular heart beats			√
Rare or Uncommon	Liver Disorder: yellowing of the skin or eyes, dark urine, abdominal pain, nausea, vomiting, loss of appetite		√	
Rare or Uncommon	Angioedema and Severe Allergic Reactions: swelling of the face, eyes, or tongue, difficulty swallowing, wheezing, hives and generalized itching, rash, fever, abdominal cramps, chest discomfort or tightness, difficulty breathing, unconsciousness			√

This is not a complete list of side effects. For any unexpected effects while taking LUPIN-TOLVAPTAN contact your doctor, nurse or pharmacist.

HOW TO STORE IT

Store LUPIN-TOLVAPTAN between 15°C to 30°C.
Keep LUPIN-TOLVAPTAN out of reach and sight of children.

Reporting Side Effects

You can report any suspected side effects associated with the use of health products to Health Canada by:

- Visiting the Web page on Adverse Reaction Reporting (canada.ca/drug-device-reporting) for information on how to report online, by mail or by fax; or
- Calling toll-free at 1-866-234-2345.

NOTE: Contact your health professional if you need information about how to manage your side effects. The Canada Vigilance Program does not provide medical advice.

MORE INFORMATION

If you want more information about LUPIN-TOLVAPTAN:

- Talk to your healthcare professional
- Find the full product monograph that is prepared for healthcare professionals and includes this Patient Medication Information by visiting the Health Canada Drug Product Database website: (<https://www.canada.ca/en/health-canada/services/drugs-health-products/drug-products/drug-product-database.html>); the manufacturer's website www.lupinpharma.ca or by calling 1-844-587-4623.

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