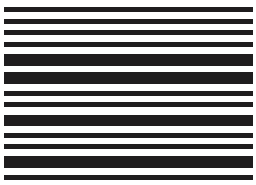


Lithium Carbonate Capsules USP

10543



2071119

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HIGHLIGHTS OF PRESCRIBING INFORMATION
These highlights do not include all the information needed to use LITHIUM CARBONATE CAPSULES safely and effectively. See full prescribing information for LITHIUM CARBONATE CAPSULES.

LITHIUM CARBONATE capsules, for oral use
Initial U.S. Approval: 1970

WARNING: LITHIUM TOXICITY

See full prescribing information for complete boxed warning.

Lithium toxicity is closely related to serum lithium concentrations, and can occur at doses close to therapeutic concentrations. Facilities for prompt and accurate serum lithium determinations should be available before initiating therapy (2.3, 5.1).

INDICATIONS AND USAGE

- Lithium is a mood-stabilizing agent indicated as monotherapy for the treatment of bipolar I disorder.
Treatment of acute manic and mixed episodes in patients 7 years and older (1)
Maintenance treatment in patients 7 years and older (1)

DOSE AND ADMINISTRATION

- Recommended starting dosage for adults and pediatric patients over 30 kg (2.2):
Capsules: 300 mg, three times daily
Recommended starting dosage for pediatric patients 20 to 30 kg (2.2):
Capsules: 300 mg twice daily
Obtain serum lithium concentration assay after 3 days, drawn 12 hours after the last oral dose and regularly until patient is stabilized.
Acute Manic or Mixed Episodes (patients 7 years and older): Titrate to serum lithium concentrations 0.8 to 1.2 mEq/L (2.2).
Maintenance Treatment for Bipolar I Disorder (patients 7 years and older): Titrate to serum lithium concentrations 0.8 to 1.2 mEq/L (2.2).

DOSE FORMS AND STRENGTHS

- Capsules: 150 mg, 300 mg, 600 mg of lithium carbonate (3)

CONTRAINDICATIONS

Known hypersensitivity to any inactive ingredient in the drug product. (4)

WARNINGS AND PRECAUTIONS

- Lithium-Induced Polyuria: May develop during initiation of treatment. Increases risk of lithium toxicity. Educate patient to avoid dehydration. Monitor for lithium toxicity and metabolic acidosis. Discontinue lithium or treat with amiloride as a therapeutic agent (5.2).
Hypонатremia: Symptoms are more severe with faster-onset hyponatremia. Dehydration from protracted sweating, diarrhea, or elevated temperatures from infection increases risk of hyponatremia and lithium toxicity. Educate patients on maintaining a normal diet with salt and staying hydrated. Monitor for and treat hyponatremia and lithium toxicity, which may necessitate a temporary reduction or cessation of lithium and infusion of serum sodium (5.3).
Lithium-Induced Chronic Kidney Disease: Associated with structural changes in patients on chronic lithium therapy. Monitor kidney function during treatment with lithium (5.4).
Encephalopathic Syndrome: Increased risk in patients treated with lithium and an antipsychotic. Monitor routinely for changes to cognitive function (5.5).
Hypothyroidism and Hyperthyroidism: Monitor thyroid function regularly (5.7).
Hypercalcemia and Hyperparathyroidism: Associated with long-term lithium use. Monitor serum calcium (5.8).

ADVERSE REACTIONS

- Adult Patients: fine hand tremor, polyuria, mild thirst, nausea, general discomfort during initial treatment (6)
Pediatric Patients (7 to 17 years): nausea/vomiting, polyuria, thyroid abnormalities, tremor, thirst/polydipsia, dizziness, rash/dermatitis, ataxia/gait disturbance, decreased appetite, and blurry vision (6)

To report SUSPECTED ADVERSE REACTIONS, contact Hetero Labs Limited at 1-866-495-1995 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DRUG INTERACTIONS

- Diuretics, NSAID, renin-angiotensin system antagonists, or metronidazole may increase lithium serum concentrations. Recommend frequent monitoring of serum lithium concentration and adjust dosage when necessary. (2.3, 7.1)
Serotonergic Agents: Increased risk of serotonin syndrome when co-administered with lithium. (5.6, 7.1)
Antipsychotics: There have been reports of neurologic adverse reactions in patients treated with lithium and an antipsychotic, ranging from extrapyramidal symptoms to neuroleptic malignant syndrome. (5.5, 7.1)

USE IN SPECIFIC POPULATIONS

- Pregnancy: May cause fetal and/or neonatal harm (8.1)
Renal Impairment: Use caution during dose selection, starting with dosages less than those for patients with normal renal function while carefully monitoring for side effects (2.5, 6)

See 17 for PATIENT COUNSELING INFORMATION and Medication Guide

Revised: 11/2022

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FULL PRESCRIBING INFORMATION

WARNING: LITHIUM TOXICITY

Lithium toxicity is closely related to serum lithium concentrations, and can occur at doses close to therapeutic concentrations. Facilities for prompt and accurate serum lithium determinations should be available before initiating treatment (See Dosage and Administration (2.3), Warnings and Precautions (5.1)).

1 INDICATIONS AND USAGE

- Lithium is a mood-stabilizing agent indicated as monotherapy for the treatment of bipolar I disorder.
Pre-treatment Screening: Evaluate renal function, vital signs, electrolytes, thyroid function should be evaluated. Concurrent medications should be assessed, and if the patient is a woman of childbearing potential, pregnancy status and potential should be considered.

2 DOSAGE AND ADMINISTRATION

- 2.1 Pre-treatment Screening
Before initiating treatment with lithium, renal function, vital signs, serum electrolytes, and thyroid function should be evaluated. Concurrent medications should be assessed, and if the patient is a woman of childbearing potential, pregnancy status and potential should be considered.

2.2 Recommended Dosage

See Table 1 for dosage recommendations for acute and maintenance treatment of bipolar I disorder in adult and pediatric patients (7 to 17 years).

Obtain serum lithium concentration assay after 3 days, drawn 12 hours after the last oral dose and regularly until patient is stabilized. Fine hand tremor, polyuria, and thirst may occur during initial therapy for the acute manic phase and may persist throughout treatment. Nausea and general discomfort may also appear during the first few days of lithium administration. These adverse reactions may subside with continued treatment, concomitant administration with food, or temporary reduction or cessation of dosage.

Table 1. Lithium Dosing for Bipolar I Disorder

Table with 8 columns: Patient Group, Formulation, Starting Dose, Dose Titration, Acute Goal (Serum Level, Usual Dose), Maintenance Goal (Serum Level, Usual Dose). Rows include Adult and Pediatric Patients over 30 kg, and Pediatric Patients 20 to 30 kg.

2.3 Serum Lithium Monitoring

Blood samples for serum lithium determination should be drawn immediately prior to the next dose when lithium concentrations are relatively stable (i.e., 12 hours after the previous dose). Total reliance must not be placed on serum concentrations alone. Accurate patient evaluation requires both clinical and laboratory analysis.

In addition to regular monitoring of serum lithium concentrations for patients on maintenance treatment, serum lithium concentrations should be monitored after any change in dosage, concurrent medication (e.g., diuretics, non-steroidal anti-inflammatory drugs, renin-angiotensin system antagonists, or metronidazole), marked increase or decrease in routinely performed strenuous physical activity (such as an exercise program) and in the event of a concomitant disease (See Boxed Warning, Warnings and Precautions (5.1), Drug Interactions (7.1)).

Patients abnormally sensitive to lithium may exhibit toxic signs at serum concentrations that are within what is considered the therapeutic range. Geriatric patients often respond to reduced dosage, and may exhibit signs of toxicity at serum concentrations ordinarily tolerated by other patients (See Specific Populations (8.5)).

2.4 Dosage Adjustments during Pregnancy and the Postpartum Period

If the decision is made to continue lithium treatment during pregnancy, monitor serum lithium concentrations and adjust the dosage as needed in a pregnant woman because renal lithium clearance increases during pregnancy. Avoid sodium restriction or diuretic administration. To decrease the risk of postpartum lithium intoxication, decrease or discontinue lithium therapy two to three days before the expected delivery date to reduce neonatal concentrations and reduce the risk of maternal lithium intoxication due to the change in vascular volume which occurs during delivery. At delivery, vascular volume rapidly decreases and the renal clearance of lithium may decrease to pre-pregnancy concentrations. Restart treatment at the preconception dose when the patient is medically stable after delivery with careful monitoring of serum lithium concentrations (See Warnings and Precautions (5.1) and Use in Specific Populations (8.1)).

2.5 Dosage Adjustments for Patients with Renal Impairment

Start patients with mild to moderately impaired renal function (creatinine clearance 30 to 89 mL/min evaluated by Cockcroft-Gault) with dosages less than those for patients with normal renal function (See Dosage and Administration (2.2)). Titrate slowly while frequently monitoring serum lithium concentrations and monitoring for signs of lithium toxicity. Lithium is not recommended for use in patients with severe renal impairment (creatinine clearance less than 30 mL/min evaluated by Cockcroft-Gault) (See Use in Specific Populations (8.6)).

3 DOSAGE FORMS AND STRENGTHS

Each 150 mg capsule for oral administration contains: lithium carbonate, USP 150 mg and is a white/white size '4' hard gelatin capsules, imprinted with '97' on body and 'H' on cap, containing white to off-white powder.

Each 300 mg capsule for oral administration contains: lithium carbonate, USP 300 mg and is a pink/pink size '1' hard gelatin capsules, imprinted with '98' on body and 'H' on cap, containing white to off-white powder.

Each 600 mg capsule for oral administration contains: lithium carbonate, USP 600 mg and is a pink/white size '0EL' hard gelatin capsules, imprinted with '141' on body and 'H' on cap, containing white to off-white powder.

4 CONTRAINDICATIONS

Lithium is contraindicated in patients with known hypersensitivity to any inactive ingredient in the lithium carbonate capsule or lithium citrate products (See Adverse Reactions (6)).

5 WARNINGS AND PRECAUTIONS

5.1 Lithium Toxicity

The toxic concentrations for lithium (>1.5 mEq/L) are close to the therapeutic range (0.8 to 1.2mEq/L). Some patients abnormally sensitive to lithium may exhibit toxic signs at serum concentrations that are considered within the therapeutic range (See Boxed Warning, Dosage and Administration (2.3)). Lithium may take up to 24 hours to distribute into brain tissue, so occurrence of acute toxicity symptoms may be delayed.

Neurological signs of lithium toxicity range from mild neurological adverse reactions such as fine tremor, lightheadedness, lack of coordination, and weakness; to moderate manifestations like giddiness, apathy, drowsiness, hyperreflexia, muscle twitching, ataxia, blurred vision, tinnitus, and slurred speech; and severe manifestations such as clonus, confusion, seizure, coma, and death. In rare cases, neurological sequelae may persist despite discontinuing lithium treatment and may be associated with cerebellar atrophy. Cardiac manifestations involve electrocardiographic changes, such as prolonged QT interval, ST and T-wave changes and myocarditis. Renal manifestations include urine concentrating defect, nephrogenic diabetes insipidus, and renal failure. Respiratory manifestations include dyspnea, aspiration pneumonia, and respiratory failure. Gastrointestinal manifestations include nausea, vomiting, diarrhea, and bloating. No specific antidote for lithium poisoning is known (See Overdosage (10)).

The risk of lithium toxicity is increased by:

- Recent onset of concurrent febrile illness
Concomitant administration of drugs which increase lithium serum concentrations by pharmacokinetic interactions or drugs affecting kidney function (See Drug Interactions (7)).
Acute ingestion
Impaired renal function
Volume depletion or dehydration
Significant cardiovascular disease
Changes in electrolyte concentrations (especially sodium and potassium)

Monitor for signs and symptoms of lithium toxicity. If symptoms occur, decrease dosage or discontinue lithium treatment.

5.2 Lithium-Induced Polyuria

Chronic lithium treatment may be associated with diminution of renal concentrating ability, occasionally presenting as nephrogenic diabetes insipidus, with polyuria and polydipsia. The concentrating defect and natriuretic effect characteristic of this condition may develop within weeks of lithium initiation. Lithium can also cause renal tubular acidosis, resulting in hyperchloremic metabolic acidosis. Such patients should be carefully managed to avoid dehydration with resulting lithium retention and toxicity. This condition is usually reversible when lithium is discontinued, although for patients treated with long-term lithium, nephrogenic diabetes insipidus may be only partly reversible upon discontinuation of lithium. Amiloride may be considered as a therapeutic agent for lithium-induced nephrogenic diabetes insipidus.

5.3 Hyponatremia

Lithium can cause hyponatremia by decreasing sodium reabsorption by the renal tubules, leading to sodium depletion. Therefore, it is essential for patients receiving lithium treatment to maintain a normal diet, including salt, and an adequate fluid intake (2,500 to 3,000 mL) at least during the initial stabilization period. Decreased tolerance to lithium has also been reported to ensue from protracted sweating or diarrhea and, if such occur, supplemental fluid and salt should be administered under careful medical supervision and lithium intake reduced or suspended until the condition is resolved. In addition, concomitant infection with elevated temperatures may also necessitate a temporary reduction or cessation of medication.

- reuptake inhibitors (SNRIs) (SSRIs)
tricyclic antidepressants
antipsychotic medicines
tryptophan
St John's Wort
fenfluramine
buspirone

Your healthcare provider can tell you if it is safe to take lithium carbonate with your other medicines. Do not start or stop any medicines while taking lithium carbonate without talking to your healthcare provider first.

Know the medicines you take. Keep a list of your medicines to show your healthcare provider and pharmacist when you get a new medicine.

How should I take lithium carbonate?

- Take your lithium carbonate exactly as prescribed by your healthcare provider.
Your healthcare provider will do certain blood tests before starting and during treatment with lithium carbonate.
Your healthcare provider may change your dose if needed. Do not change your dose on your own.
Do not double your dose if a dose is missed. Talk with your healthcare provider if you miss a dose.
Do not stop taking lithium carbonate suddenly without talking to your healthcare provider.
Your healthcare provider may change your lithium carbonate dose to make sure you are taking the dose that is right for you.
If you take too much lithium carbonate, call your healthcare provider or poison control center, or go to the nearest hospital emergency room right away, in case of poisoning, call your poison control center at 1-800-222-1222.

What should I avoid while taking lithium carbonate?

- Do not drink, operate heavy machinery, or do other dangerous activities when you start taking lithium carbonate, when your dose is changed, or until you know how lithium carbonate affects you.
Lithium carbonate can make you sleepy. Talk to your healthcare provider about these activities.
Avoid becoming overheated or dehydrated during exercise and in hot weather. Follow your healthcare provider instructions about the type and amount of liquids you should drink. In some cases, drinking too much liquid can be as unsafe as not drinking enough.
Do not change the amount of salt in your diet.
Changing the amount of salt in your diet could change the amount of lithium carbonate in your blood.

What are the possible side effects of lithium carbonate?

See "What is the most important information I should know about lithium carbonate?"
Lithium carbonate may cause serious side effects, including:
kidney problems. People who take lithium carbonate may have to urinate often (polyuria) and have other kidney problems that may affect how their kidneys work. These problems can happen within a few weeks of starting to take lithium carbonate or after taking lithium carbonate for a long time.
low levels of sodium (salt) in your blood (hyponatremia). Lithium carbonate can cause you to lose sodium. Talk to your healthcare provider about your diet and how much fluid you are drinking when starting lithium carbonate. If you have been sweating more than usual or have had diarrhea, you may need extra salt and more fluids. Talk to your healthcare provider if this happens.
neurological problems. People who take lithium

MEDICATION GUIDE
Lithium Carbonate (lith ee um kar' bo nate) Capsules, USP
What is the most important information I should know about lithium carbonate?
Lithium carbonate can cause serious side effects, including:
Too much lithium in your blood (lithium toxicity). Lithium toxicity that can cause death may happen even if the lithium level in your blood is close to the right level for you. Your healthcare provider will need to monitor your blood levels of lithium to find the best dose for you. Take your lithium carbonate exactly as your healthcare provider tells you to take it. Stop taking lithium carbonate and call your healthcare provider right away if you have any symptoms of lithium toxicity including:
abnormal heartbeat
vomiting
drowsiness
diarrhea
weak muscles
blurred vision
clumsiness
ringing in your ears
muscle twitching

Other symptoms may include:

- lightheadedness
confusion
bloating
mood changes
slurred speech
breathing problems
seizure
coma

What is lithium carbonate?

Lithium carbonate is prescription medicine called mood-stabilizing agent used alone (monotherapy) for the acute (short-term) treatment of people 7 years of age and older with manic and mixed episodes that happen with bipolar I disorder.
maintenance treatment of bipolar I disorder in people 7 years of age and older.
It is not known if lithium carbonate is safe and effective in children under 7 years of age with bipolar I disorder.

Who should not take lithium carbonate?

Do not take lithium carbonate if you are allergic to lithium or any of the ingredients in lithium carbonate capsules. See the end of this Medication Guide for a complete list of ingredients in lithium carbonate capsules.
What should I tell my healthcare provider before taking lithium carbonate?
Before taking lithium carbonate, tell your healthcare provider if you:
have kidney problems
have heart problems
have breathing problems
have thyroid problems
are pregnant or plan to become pregnant. Lithium carbonate may harm your unborn baby
are breastfeeding or plan to breastfeed. Lithium carbonate can pass into your breastmilk and may harm your baby. You should not breastfeed during treatment with lithium carbonate. Talk to your healthcare provider about the best way to feed your baby if you take lithium carbonate.

Tell your healthcare provider about all the medicines you take, including prescription, over-the-counter medicines, vitamins, and herbal supplements.

Using lithium carbonate with certain other medicines may affect each other causing possible side effects. Lithium carbonate may affect the way other medicines work, and other medicines may affect how lithium carbonate works.

Especially tell your healthcare provider if you take:

- MAOIs
serotonin reuptake inhibitors
serotonin norepinephrine reuptake inhibitors

Table with 2 columns: Property and Value. Rows include Dimensions (300 x 550 mm), Country (USA), Spec (Bible Paper 40 GSM), and Pantone Colours (Black).



System Organ Class/ Preferred Term	Placebo N=28 ^a	Lithium N=53 ^b
Thirst/polydipsia	11	28
Decreased appetite	4	9
Nervous system disorders		
Ataxia/gait disturbance	0	13
Blurry vision	0	9
Disorientation	0	6
Dizziness	7	23
Tremor	7	32
Skin and subcutaneous tissue disorders		
Rash/dermatitis	0	13

Adult Patients:

The following adverse reactions have been identified following use of lithium. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

Central Nervous System: tremor, muscle hyperirritability (fasciculations, twitching, clonic movements of whole limbs), hypertonicity, ataxia, choreoathetotic movements, hyperactive deep tendon reflexes, extrapyramidal symptoms including acute dystonia, cogwheel rigidity, blackout spells, epileptiform seizures, slurred speech, dizziness, vertigo, downbeat nystagmus, incontinence of urine or feces, somnolence, psychomotor retardation, restlessness, confusion, stupor, coma, tongue movements, tics, tinnitis, hallucinations, poor memory, slowed intellectual functioning, started response, worsening of organic brain syndromes, myasthenic syndromes (rarely).

EEG Changes: diffuse slowing, widening of frequency spectrum, potentiation and disorganization of background rhythm.

Cardiovascular: conduction disturbance (mostly sinus node dysfunction with possibly severe sinus bradycardia and sinoatrial block), ventricular tachyarrhythmia, peripheral vasculopathy (resembling Raynaud's Syndrome).

ECG Changes: reversible flattening, isoelectricity or rarely inversion of T-waves, prolongation of the QTc interval.

Gastrointestinal: anorexia, nausea, vomiting, diarrhea, gastritis, salivary gland swelling, abdominal pain, excessive salivation, flatulence, indigestion.

Genitourinary: glycosuria, decreased creatinine clearance, albuminuria, oliguria, and symptoms of nephrogenic diabetes insipidus including polyuria, thirst, and polydipsia.

Dermatologic: drying and thinning of hair, alopecia, anesthesia of skin, chronic folliculitis, xerosis cutis, psoriasis onset or exacerbation, generalized pruritus with or without rash, cutaneous ulcers, angioedema, drug reaction with eosinophilia and systemic symptoms (DRESS).

Autonomic Nervous System: blurred vision, dry mouth, impotence/sexual dysfunction.

Miscellaneous: fatigue, lethargy, transient scotoma, exophthalmos, dehydration, weight loss, leukocytosis, headache, resistant hyperglycemia, hypermagnesemia, excessive weight gain, edematous swelling of ankles or wrists, dysgeusia/taste distortion (e.g., metallic or salty taste), thirst, swollen lips, tightness in chest, swollen and/or painful joints, fever, polyarthralgia, and dental caries.

7 DRUG INTERACTIONS**7.1 Drugs Having Clinically Important Interactions with Lithium****Table 4: Clinically Important Drug Interactions with Lithium**

Diuretics	
<i>Clinical Impact:</i>	Diuretic-induced sodium loss may reduce lithium clearance and increase serum lithium concentrations.
<i>Intervention:</i>	More frequent monitoring of serum electrolyte and lithium concentrations. Reduce lithium dosage based on serum lithium concentration and clinical response [see <i>Dosage and Administration</i> (2.3), <i>Warnings and Precautions</i> (5.3)].
Non-Steroidal Anti-inflammatory Drugs (NSAID)	
<i>Clinical Impact:</i>	NSAID decrease renal blood flow, resulting in decreased renal clearance and increased serum lithium concentrations.
<i>Intervention:</i>	More frequent serum lithium concentration monitoring. Reduce lithium dosage based on serum lithium concentration and clinical response [see <i>Dosage and Administration</i> (2.3)].
Renin-Angiotensin System Antagonists	
<i>Clinical Impact:</i>	Concomitant use increase steady-state serum lithium concentrations.
<i>Intervention:</i>	More frequent monitoring of serum lithium concentration. Reduce lithium dosage based on serum lithium concentration and clinical response [see <i>Dosage and Administration</i> (2.3)].
Serotonergic Drugs	
<i>Clinical Impact:</i>	Concomitant use can precipitate serotonin syndrome.
<i>Intervention:</i>	Monitor patients for signs and symptoms of serotonin syndrome, particularly during lithium initiation. If serotonin syndrome occurs, consider discontinuation of lithium and/or concomitant serotonergic drugs [see <i>Warnings and Precautions</i> (5.6)].
Nitroimidazole Antibiotics	
<i>Clinical Impact:</i>	Concomitant use may increase serum lithium concentrations due to reduced renal clearance.
<i>Intervention:</i>	More frequent monitoring of serum lithium concentration. Reduce lithium dosage based on serum lithium concentration and clinical response [see <i>Dosage and Administration</i> (2.3)].
Acetazolamide, Urea, Xanthine Preparations, Alkalinizing Agents	
<i>Clinical Impact:</i>	Concomitant use can lower serum lithium concentrations by increasing urinary lithium excretion.
<i>Intervention:</i>	More frequent serum lithium concentration monitoring. Increase lithium dosage based on serum lithium concentration and clinical response [see <i>Dosage and Administration</i> (2.3)].
Methyldopa, Phenytoin and Carbamazepine	
<i>Clinical Impact:</i>	Concomitant use may increase risk of adverse reactions of these drugs.
<i>Intervention:</i>	Monitor patients closely for adverse reactions of methyldopa, phenytoin, and carbamazepine.
Iodide Preparations	
<i>Clinical Impact:</i>	Concomitant use may produce hypothyroidism.
<i>Intervention:</i>	Monitor patients for signs or symptoms of hypothyroidism [see <i>Warnings and Precautions</i> (5.7)].
Calcium Channel Blocking Agents (CCB)	
<i>Clinical Impact:</i>	Concomitant use may increase the risk of neurologic adverse reactions in the form of ataxia, tremors, nausea, vomiting, diarrhea and/or tinnitus.
<i>Intervention:</i>	Monitor for neurologic adverse reactions.
Atypical and Typical Antipsychotic Drugs	
<i>Clinical Impact:</i>	Reports of neurotoxic reactions in patients treated with both lithium and an antipsychotic, ranging from extrapyramidal syndromes to neuroleptic malignant syndrome, as well as reports of an encephalopathic syndrome in few patients treated with concomitant therapy [see <i>Warnings and Precautions</i> (5.5)].
<i>Intervention:</i>	Monitor for neurologic adverse reactions.
Sodium-Glucose Cotransporter 2 (SGLT2) Inhibitor	
<i>Clinical Impact:</i>	Concomitant use of lithium with an SGLT2 inhibitor may decrease serum lithium concentrations.
<i>Intervention:</i>	Monitor serum lithium concentration more frequently during SGLT2 inhibitor initiation and dosage changes.
Neuromuscular Blocking Agents	
<i>Clinical Impact:</i>	Lithium may prolong the effects of neuromuscular blocking agents.
<i>Intervention:</i>	Monitor for prolonged paralysis.

8 USE IN SPECIFIC POPULATIONS**8.1 Pregnancy****Risk Summary:**

Lithium may cause harm when administered to a pregnant woman. Early voluntary reports to international birth registries suggested an increase in cardiovascular malformations, especially for Ebstein's anomaly, with first trimester use of lithium. Subsequent case-control and cohort studies indicate that the increased risk for cardiac

malformations is likely to be small; however, the data are insufficient to establish a drug-associated risk. There are concerns for maternal and/or neonatal lithium toxicity during late pregnancy and the postpartum period [see *Clinical Considerations*]. Published animal developmental and toxicity studies in mice and rats report an increased incidence of fetal mortality, decreased fetal weight, increased fetal skeletal abnormalities, and cleft palate (mouse fetuses only) with oral doses of lithium that produced serum concentrations similar to the human therapeutic range. Other published animal studies report adverse effects on embryonic implantation in rats after lithium administration. Advise pregnant women of the potential risk to a fetus.

The background risk of major birth defects and miscarriage for the indicated population(s) is unknown. In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2 to 4% and 15 to 20%, respectively.

Clinical Considerations:
Dose Adjustments During Pregnancy and the Postpartum Period: If the decision is made to continue lithium treatment during pregnancy, serum lithium concentrations should be monitored and the dosage adjusted during pregnancy. Two to three days prior to delivery, lithium dosage should be decreased or discontinued to reduce the risk of maternal and/or neonatal toxicity. Lithium may be restarted in the post-partum period at preconception doses in medically stable patients as long as serum lithium levels are closely monitored [see *Dosage and Administration* (2.4), *Warnings and Precautions* (5.1)].

Fetal/Neonatal Adverse Reactions: Lithium toxicity may occur in neonates who were exposed to lithium in late pregnancy. A floppy baby syndrome including neurological, cardiac, and hepatic abnormalities that are similar to those seen with lithium toxicity in adults have been observed. Symptoms include hypotonia, respiratory distress syndrome, cyanosis, lethargy, feeding difficulties, depressed neonatal reflexes, neonatal depression, apnea, and bradycardia. Monitor neonates and provide supportive care until lithium is excreted and toxic signs disappear, which may take up to 14 days.

Consider fetal echocardiography between 16 and 20 weeks gestation in a woman with first trimester lithium exposure because of the potential increased risk of cardiac malformations.

8.2 Lactation

Risk Summary:
Limited published data reports the presence of lithium carbonate in human milk with breast milk levels measured at 0.12 to 0.7 mEq or 40 to 45% of maternal plasma levels. Infants exposed to lithium during breastfeeding may have plasma levels that are 30 to 40% of maternal plasma levels. Signs and symptoms of lithium toxicity such as hypertonia, hyperthermia, cyanosis, and ECG changes have been reported in some breastfed neonates and infants. Increased prolactin levels have been measured in lactating women, but the effects on milk production are not known. Breastfeeding is not recommended with maternal lithium use; however, if a woman chooses to breastfeed, the infant should be closely monitored for signs of lithium toxicity. Discontinue breastfeeding if a breastfed infant develops lithium toxicity.

Clinical Considerations:
Consider regular monitoring of lithium levels and thyroid function in a breastfed infant.

8.4 Pediatric Use

The safety and effectiveness of lithium for monotherapy treatment of acute manic or mixed episodes of bipolar I disorder and maintenance monotherapy of bipolar I disorder in pediatric patients ages 7 to 17 years of age have been established in an acute-phase clinical trial of 8 weeks in duration followed by a 28-week randomized withdrawal phase [see *Dosage and Administration* (2.1), *Adverse Reactions* (6.1), *Clinical Pharmacology* (12.3), *Clinical Studies* (14)].

The safety and effectiveness of lithium has not been established in pediatric patients less than 7 years of age with bipolar I disorder.

8.5 Geriatric Use

Clinical studies of lithium carbonate tablets did not include sufficient numbers of subjects aged 65 and over to determine whether they respond differently from younger subjects. Other reported clinical experience has not identified differences in response between the elderly and younger patients. In general, dose selection for an elderly patient should be cautious, usually starting at the low end of the dosing range, reflecting the greater frequency of decreased hepatic, renal, or cardiac function, and of concomitant disease or other treatment.

Lithium is known to be substantially excreted by the kidneys, and the risk of toxic reactions to this drug may be greater in patients with impaired renal function. Because elderly patients are more likely to have decreased renal function, care should be taken in dose selection, and it may be useful to monitor renal function.

8.6 Renal Impairment

As lithium is eliminated primarily through the kidney, lithium renal clearance is decreased in patients with abnormal renal function, and the risk of lithium intoxication increases considerably in this setting. Lithium should not be used in severe renal insufficiency (creatinine clearance less than 30 mL/min evaluated by Cockcroft-Gault), especially if the condition requires adherence to a low-sodium diet [see *Dosage and Administration* (2.5)].

Start patients with mild to moderately impaired renal function (creatinine clearance 30 to 89 mL/min evaluated by Cockcroft-Gault) with lower doses of lithium and titrate slowly while frequently monitoring serum lithium concentrations and for signs of lithium toxicity [see *Dosage and Administration* (2.5)].

10 OVERDOSAGE

The toxic concentrations for lithium (≥ 1.5 mEq/L) are close to the therapeutic concentrations [see *Warnings and Precautions* (5.1)]. At lithium concentrations greater than 3 mEq/L, patients may progress to seizures, coma, and irreversible brain damage.

Treatment:

For current information on the management of poisoning or overdose, contact the National Poison Control Center at 1-800-222-1222 or www.poisson.org.

No specific antidote for lithium poisoning is known. Mild symptoms of lithium toxicity can usually be treated by reduction in dose or cessation of the drug.

In severe cases of lithium poisoning, the goal of treatment is elimination of this ion from the patient. Administration of gastric lavage should be performed, but use of activated charcoal is not recommended as it does not significantly absorb lithium ions. Hemodialysis is the treatment of choice as it is an effective and rapid means of removing lithium in patients with severe toxicity. As an alternative option, urea, mannitol and aminophylline can induce a significant increase in lithium excretion. Appropriate supportive care for the patient should be undertaken. Patients with impaired consciousness should have their airway protected and it is critical to correct any volume depletion or electrolyte imbalance. Patients should be monitored to prevent hypernatremia while receiving normal saline and careful regulation of kidney function is of utmost importance.

Serum lithium concentrations should be closely monitored as there may be a rebound in serum lithium concentrations as a result of delayed diffusion from the body tissues. Likewise, during the late recovery phase, lithium should be re-administered with caution taking into account the possible release of significant lithium stores in body tissues.

11 DESCRIPTION

Each capsule for oral administration contains lithium carbonate USP, 150 mg, 300 mg or 600 mg and the following inactive ingredients: gelatin, sodium lauryl sulfate, talc, titanium dioxide and the imprinting ink contains black iron oxide E172 dye, butyl alcohol, dehydrated alcohol, isopropyl alcohol, potassium hydroxide, propylene glycol, shellac and strong ammonia solution.

Lithium is an element of the alkali-metal group with atomic number 3, atomic weight 6.94, and an emission line at 671 nm on the flame photometer.

Lithium Carbonate USP is a white, light, alkaline powder with molecular formula Li₂CO₃ and molecular weight 73.89.

12 CLINICAL PHARMACOLOGY**12.1 Mechanism of Action**

The mechanism of action of lithium as a mood stabilizing agent is unknown.

12.3 Pharmacokinetics**Absorption:**

After oral administration, lithium is reported to be completely absorbed in the upper gastrointestinal tract. Peak serum concentrations (T_{max}) occur 0.25 to 3 hours after oral administration of immediate release preparations and 2 to 6 hours after sustained-release preparations.

Distribution:

The distribution space of lithium approximates that of total body water, and the plasma protein binding is negligible. After equilibrium, the apparent volume of distribution is 0.7 to 1 L/kg.

Metabolism:

Lithium is not metabolized.

Excretion:

Lithium is primarily excreted in urine, proportionally to its serum concentration. Lithium is filtered by the glomerulus, and 80% is reabsorbed by passive diffusion in the proximal tubule. The elimination half-life of lithium is approximately 18 to 36 hours. Lithium excretion in feces is insignificant.

Specific Populations:

Pediatric Use. A pharmacokinetic study of lithium was performed in 39 subjects with bipolar I disorder. Both apparent clearance and apparent volume of distribution increase as body weight increases. A lower dose in patients < 30 kg is necessary to achieve lithium exposures in pediatric patients similar to those observed in adults treated at recommended doses of lithium [see *Dosage and Administration* (2.2)]. The estimated plasma clearance was 0.59 L/h, 0.79 L/h and 1.17 L/h for pediatric patients weighing 20 kg, 30 kg and 50 kg, respectively.

13 NONCLINICAL TOXICOLOGY**13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility**

Carcinogenesis:
There have been no long-term studies performed in animals to evaluate the carcinogenic potential of lithium.

Mutagenesis:

There have been no adequate studies conducted to evaluate the mutagenic and genotoxic potential of lithium.

Impairment of Fertility:

There have been no adequate studies performed in animals at current standards to evaluate the effect of lithium treatment on fertility. However, published studies in male mice and rats administered repeated daily dosing of lithium carbonate report adverse effects on male reproductive organs, decreased spermatogenesis and decreased testosterone levels.

14 CLINICAL STUDIES

The safety and efficacy of lithium as a treatment for acute manic or mixed episodes of bipolar I disorder in pediatric patients (ages 7 to ≤18 years) was demonstrated in an 8-week, randomized, placebo-controlled, parallel group study (NCT01166425). In this study, 81 patients with a Young Mania Rating Scale (YMRS) score of 20 or more were randomized to receive lithium or placebo in a 2:1 ratio. Patients weighing more than 30 kg started lithium at 300 mg three times daily (900 mg/day) and could increase their dose by 300 mg every 3 days. Patients weighing 20 to 30 kg started lithium at 300 mg twice daily (600 mg/day) and could increase their dose by 300 mg weekly. No patients weighing less than 20 kg were enrolled. Lithium (mean serum level 0.98 ± 0.47 mEq/L) was statistically significantly superior to placebo in decreasing acute mania or mixed states as measured by the YMRS (see Table 5).

In a 28-week randomized withdrawal analysis, 31 pediatric patients stabilized on lithium were assigned to either continue lithium or switch to placebo. The group receiving lithium demonstrated superiority to those receiving placebo in all-cause discontinuation (see Table 5).

Table 5: Primary Efficacy Results

Analysis	Treatment Group	Change From Baseline at Week 8 in YMRS Summary Score			
		N	Mean Baseline Score (SD)	LS Mean Change from Baseline (SE)	Difference ¹ (95% CI)
Acute Efficacy	Lithium:	53	29.5 (6.6)	-12.9 (3.1)	-5.5 (-10.5, -0.5)
	Placebo:	28	30.0 (6.0)	-7.3 (3.1)	
All-cause Discontinuation					
Analysis	Treatment Group ²	N	Number of Discontinued Subjects		Hazard Ratio ³ (95% CI)
			Lithium:	Placebo:	
Randomized	Lithium:	17	7	41.2%	0.28 (0.10, 0.78)
Withdrawal	Placebo:	14	11	78.6%	

SD: standard deviation; SE: standard error; LS Mean: least-squares mean; CI: confidence interval.

1. Difference (drug minus placebo) in least-squares mean change from baseline.
2. Patients analyzed by received treatment.
3. Lithium to placebo.

16 HOW SUPPLIED/STORAGE AND HANDLING**Lithium Carbonate Capsules, USP**

Lithium Carbonate Capsules USP, 150 mg are white/white size '4' hard gelatin capsules, imprinted with '97' on body and 'H' on cap, containing white to off-white powder. They are supplied in

Bottles of 30 Capsules	NDC 31722-544-30
Bottles of 100 Capsules	NDC 31722-544-01
Bottles of 500 Capsules	NDC 31722-544-05
Bottles of 1000 Capsules	NDC 31722-544-10

Lithium Carbonate Capsules USP, 300 mg are pink/pink size '1' hard gelatin capsules, imprinted with '98' on body and 'H' on cap, containing white to off-white powder. They are supplied in

Bottles of 30 Capsules	NDC 31722-545-30
Bottles of 100 Capsules	NDC 31722-545-01
Bottles of 500 Capsules	NDC 31722-545-05
Bottles of 1000 Capsules	NDC 31722-545-10
Bottles of 5000 Capsules	NDC 31722-545-50
Blister Pack of 3x10's	NDC 31722-545-03

Lithium Carbonate Capsules USP, 600 mg are pink/white size 'OEL' hard gelatin capsules, imprinted with '141' on body and 'H' on cap, containing white to off-white powder. They are supplied in

Bottles of 30 Capsules	NDC 31722-546-30
Bottles of 100 Capsules	NDC 31722-546-01
Bottles of 500 Capsules	NDC 31722-546-05
Bottles of 1000 Capsules	NDC 31722-546-10

Storage

Store at 20° to 25°C (68° to 77°F) [see USP Controlled Room Temperature]. Dispense in a tight, child-resistant container as defined in the USP/NF. PROTECT FROM MOISTURE.

17 PATIENT COUNSELING INFORMATION

Advise the patient to read FDA-approved patient labeling (Medication Guide).

Dosage and Administration:

Advise patients that lithium is a mood stabilizer, and should only be taken as directed. Emphasize the importance of compliance with the prescribed treatment and to not adjust the dose of lithium without first consulting their healthcare provider. Inform patients that they will need to have regular blood draws to determine if their dose of lithium is appropriate.

Inform patients to not double the dose if a dose is missed, due to the complexity of individualized dosing and potential for lithium toxicity [see *Dosage and Administration* (2), *Warnings and Precautions* (5.1)].

Lithium Toxicity:

Inform patients on adverse reactions related to lithium toxicity that require medical attention. Advise patients to inform their healthcare provider if clinical signs of lithium toxicity such as diarrhea, vomiting, tremor, lack of muscle coordination, drowsiness, abnormal heart rhythm or muscular weakness occur [see *Warnings and Precautions* (5.1)].

Lithium-Induced Polyuria:

Advise patients on the adverse reactions related to lithium-induced polyuria, when to seek medical attention, and the importance of maintaining normal diet with salt and staying hydrated [see *Warnings and Precautions* (5.2)].

Hypонатremia:

Counsel patients on the adverse reactions of hyponatremia, when to seek medical attention, the importance of maintaining a normal diet including adequate salt intake and staying hydrated [see *Warnings and Precautions* (5.3)]. Salt supplements and additional fluids may be required if excessive losses occur.

Serotonin Syndrome:

Counsel patients about the risk of serotonin syndrome, particularly with the concomitant use of lithium with other serotonergic drugs including SSRIs, SNRIs, triptans, tricyclic antidepressants, fenfluramine, tramadol, tryptophan, buspirone, St. John's Wort, and with drugs that impair metabolism of serotonin (in particular, MAOIs, both those intended to treat psychiatric disorders and also others, such as linezolid) [see *Warnings and Precautions* (5.6) and *Drug Interactions* (7)].

Drug Interactions:

Advise patients that many drugs can interact with lithium and to inform their doctor and pharmacist if they are taking any over the counter medication, including herbal medication, or are started on a new prescription [see *Drug Interactions* (7)].

Somnolence:

Tell patients that lithium may cause somnolence particularly when initiating treatment and to be cautious about operating vehicles or hazardous machinery, until they are reasonably certain that lithium treatment does not affect them adversely [see *Adverse Reactions* (6)].

Pregnancy:

Advise pregnant women of the potential risk to a fetus and/or neonate [see *Use in Specific Populations* (8.1)].

Lactation:

Advise women that breastfeeding is not recommended during treatment with lithium [see *Use in Specific Populations* (8.2)].



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India

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carbamate with certain other medicines called antipsychotics may have symptoms such as weakness, tiredness, fever, tremors, and confusion. Talk to your healthcare provider if this happens. Ask if you are not sure about the medicines you take.

- **serotonin syndrome.** A potentially life-threatening problem called serotonin syndrome can happen when you take lithium carbonate while you take certain medicines called serotonergic and MAOIs. Symptoms of serotonin syndrome include:
 - agitation
 - seeing things that are not there
 - confusion
 - coma
 - rapid pulse
 - high or low blood pressure
 - dizziness
 - sweating
 - flushing
 - fever
 - tremors
 - stiff muscles
 - muscle twitching
 - become unstable
 - seizures
 - nausea
 - vomiting
 - diarrhea
- **thyroid problems.**
 - high calcium levels in your blood (hypercalcemia) and changes in your parathyroid gland (hyperparathyroidism) that may not go away when you stop taking lithium carbonate.
 - **heart problems.** People who take lithium carbonate may find out they also have a heart problem called Brugada Syndrome. People who have unexplained fainting or who have a family history of sudden unexplained death before 45 years of age may have Brugada Syndrome and not know it. If you faint or feel abnormal heartbeats, talk to your healthcare provider right away.
 - **increased pressure in the brain and swelling in the eye (pseudotumor cerebri)** that can cause vision problems or blindness. If you have severe headaches behind your eyes, ringing in the ears, blurred vision, double vision, or brief periods of blindness, talk to your health care provider right away.
- The most common side effects of lithium carbonate, include:
 - Adults with manic or mixed episodes of bipolar I disorder:
 - hand trembling
 - increased urination
 - excessive thirst
 - nausea
 - general discomfort when you start treatment
 - Children 7 to 17 years of age with manic or mixed episodes of bipolar I disorder:
 - excessive urination
 - thyroid problems
 - hand trembling
 - excessive thirst
 - dizziness
 - rash
 - difficulty walking
 - decreased appetite
 - blurred vision
 - nausea
 - vomiting
- These are not all the possible side effects of lithium carbonate.

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

How should I store lithium carbonate capsules?
Store lithium carbonate capsules at room temperature, between 68° to 77°F (20° to 25°C).

Keep capsules dry and keep in a tightly closed container.

Keep lithium carbonate capsules, and all medicines out of the reach of children.