

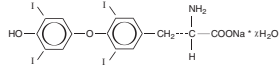
# LEVOXYL®

(levothyroxine sodium, USP)

## FOR ORAL ADMINISTRATION

### Inactive Ingredients

Microcrystalline cellulose, croscarmellose sodium and magnesium stearate. The following are the coloring additives per tablet strength:



| Strength (mcg) | Color additive(s)   |
|----------------|---|
| 25             | FD&C Yellow No. 6 Aluminum Lake   |
| 50             | None  |
| 75             | FD&C Blue No. 1 Aluminum Lake, D&C Red No. 30 Aluminum Lake                                     |
| 88             | FD&C Yellow No. 6 Aluminum Lake, FD&C Blue No. 1 Aluminum Lake, D&C Yellow No. 10 Aluminum Lake |
| 100            | FD&C Yellow No. 6 Aluminum Lake, D&C Yellow No. 10 Aluminum Lake                                |
| 112            | FD&C Yellow No. 6 Aluminum Lake, FD&C Red No. 40 Aluminum Lake, D&C Red No. 30 Aluminum Lake    |
| 125            | FD&C Red No. 40 Aluminum Lake, D&C Yellow No. 10 Aluminum Lake                                  |
| 137            | FD&C Blue No. 1 Aluminum Lake   |
| 150            | FD&C Blue No. 1 Aluminum Lake, D&C Red No. 30 Aluminum Lake                                     |
| 175            | FD&C Blue No. 1 Aluminum Lake, D&C Yellow No. 10 Aluminum Lake                                  |
| 200            | D&C Red No. 30 Aluminum Lake, D&C Yellow No. 10 Aluminum Lake                                   |
| 300            | FD&C Yellow No. 6 Aluminum Lake, FD&C Blue No. 1 Aluminum Lake, D&C Yellow No. 10 Aluminum Lake |

### CLINICAL PHARMACOLOGY

Thyroid hormone synthesis and secretion is regulated by the hypothalamic-pituitary-thyroid axis. Thyrotropin-releasing hormone (TRH) from the hypothalamus stimulates secretion of thyroid-stimulating hormone (TSH) from the anterior pituitary. TSH, in turn, is the physiologic stimulus for the synthesis and secretion of thyroid hormones. L-thyronine (T<sub>1</sub>) and L-triiodothyronine (T<sub>3</sub>), by the thyroid gland. Circulating serum T<sub>3</sub> and T<sub>4</sub> levels exert a feedback effect on both TRH and TSH secretion. When serum T<sub>3</sub> and T<sub>4</sub> levels increase, TRH and TSH secretion decrease. When thyroid hormone levels decrease, TRH and TSH secretion increase.

The mechanisms by which thyroid hormones exert their physiologic actions are not completely understood, but it is thought that their principal effects are exerted through control of DNA transcription and protein synthesis. T<sub>3</sub> and T<sub>4</sub> diffuse into the cell nucleus and bind to thyroid receptor proteins attached to DNA. This hormone nuclear receptor complex activates gene transcription and synthesis of messenger RNA and cytoplasmic proteins.

Thyroid hormones regulate multiple metabolic processes and play an essential role in normal growth and development, and normal maturation of the central nervous system and bone. The metabolic actions of thyroid hormones include augmentation of cellular respiration and thermogenesis, as well as metabolism of proteins, carbohydrates and lipids. The protein anabolic effects of thyroid hormones are essential to normal growth and development.

The physiologic actions of thyroid hormones are produced predominantly by T<sub>3</sub>, the majority of which (approximately 80%) is derived from T<sub>4</sub> by deiodination in peripheral tissues.

Levothyroxine, at doses individualized according to patient response, is effective as replacement or supplementary therapy in hypothyroidism of any etiology, except transient hypothyroidism during the recovery phase of subacute thyroiditis.

Levothyroxine is also effective in the suppression of pituitary TSH secretion in the treatment or prevention of various types of euthyroid goiters, including thyroid nodules, Hashimoto's thyroiditis, multinodular goiter and, as adjunctive therapy in the management of thyrotropin-dependent well-differentiated thyroid cancer (see **INDICATIONS AND USAGE, PRECAUTIONS, DOSAGE AND ADMINISTRATION**).

### Pharmacokinetics

**Absorption** – Absorption of orally administered T<sub>4</sub> from the gastrointestinal (GI) tract ranges from 40% to 80%. The majority of the levothyroxine dose is absorbed from the jejunum and upper ileum. The relative bioavailability of LEVOXYL® tablets, compared to an equal nominal dose of oral levothyroxine sodium solution, is approximately 98%. T<sub>4</sub> absorption is increased by fasting, and decreased in malabsorption syndromes and by certain foods such as soybean infant formula. Dietary fiber decreases bioavailability of T<sub>4</sub>. Absorption may also decrease with age. In addition, many drugs and foods affect T<sub>4</sub> absorption (see **PRECAUTIONS, Drug Interactions and Drug-Food Interactions**).

**Distribution** – Circulating thyroid hormones are greater than 99% bound to plasma proteins, including thyroxine-binding globulin (TBG), thyroxine-binding prealbumin (TBPA), and albumin (TBA), whose capacities and affinities vary for each hormone. The higher affinity of both TBG and TBPA for T<sub>4</sub> partially explains the higher serum levels, slower metabolic clearance, and longer half-life of T<sub>4</sub> compared to T<sub>3</sub>. Protein-bound thyroid hormones exist in reverse equilibrium with small amounts of free hormone. Only unbound hormone is metabolically active. Many drugs and physiologic conditions affect the binding of thyroid hormones to serum proteins (see **PRECAUTIONS, Drug Interactions and Drug-Laboratory Test Interactions**). Thyroid hormones do not readily cross the placental barrier (see **PRECAUTIONS, Pregnancy**).

**Metabolism** – T<sub>4</sub> is slowly eliminated (see **TABLE 1**). The major pathway of thyroid hormone metabolism is through sequential deiodination. Approximately eighty-percent of circulating T<sub>4</sub> is derived from peripheral T<sub>4</sub> by monodeiodination. The liver is the major site of degradation for both T<sub>4</sub> and T<sub>3</sub>, with T<sub>4</sub> deiodination also occurring at a number of additional sites, including the kidney and other tissues. Approximately 80% of the daily dose of T<sub>4</sub> is deiodinated to yield equal amounts of T<sub>3</sub> and reverse T<sub>3</sub> (rT<sub>3</sub>). T<sub>3</sub> and rT<sub>3</sub> are further deiodinated to diiodothyronine. Thyroid hormones are also metabolized via conjugation with glucuronides and sulfates and excreted directly into the bile and gut where they undergo enterohepatic recirculation.

**Elimination** – Thyroid hormones are primarily eliminated by the kidneys. A portion of the conjugated hormone reaches the colon unchanged and is eliminated in the feces. Approximately 20% of T<sub>4</sub> is eliminated in the stool. Urinary excretion of T<sub>4</sub> decreases with age.

| Table 1: Pharmacokinetic Parameters of Thyroid Hormones in Euthyroid Patients |                        |                  |                       |                                  |  |
|---|------------------------|------------------|-----------------------|----------------------------------|--|
| Hormone   | Ratio in Thyroglobulin | Biologic Potency | 1/2 <sup>(mean)</sup> | Protein Binding (%) <sup>1</sup> |  |
| Levothyroxine (T <sub>4</sub> )   | 10–20                  | 1                | 6–7 <sup>2</sup>      | 99.96                            |  |
| Liothyronine (T <sub>3</sub> )  | 1                      | 4                | 2                     | 99.5                             |  |

<sup>1</sup> 3 to 4 days in hyperthyroidism; 9 to 10 days in hypothyroidism;  
<sup>2</sup> Includes TBG, TBPA, and TBA

### INDICATIONS AND USAGE

Levothyroxine sodium is used for the following indications:

**Hypothyroidism** – As replacement or supplemental therapy in congenital or acquired hypothyroidism of any etiology, except transient hypothyroidism during the recovery phase of subacute thyroiditis. Specific indications include: primary (thyroidal), secondary (pituitary), and tertiary (hypothalamic) hypothyroidism and subclinical hypothyroidism. Primary hypothyroidism may result from functional deficiency, primary atrophy, partial or total congenital absence of the thyroid gland, or from the effects of surgery, radiation, or drugs, with or without the presence of goiter.

**Pituitary TSH Suppression** – In the treatment or prevention of various types of euthyroid goiters (see **WARNINGS and PRECAUTIONS**), including thyroid nodules (see **WARNINGS and PRECAUTIONS**), subacute or chronic lymphocytic thyroiditis (Hashimoto's thyroiditis), multinodular goiter (see **WARNINGS and PRECAUTIONS**) and, as an adjunct to surgery and radioiodine therapy in the management of thyrotropin-dependent well-differentiated thyroid cancer.

### CONTRAINDICATIONS

Levothyroxine is contraindicated in patients with untreated subclinical (suppressed serum TSH level with normal T<sub>3</sub> and T<sub>4</sub> levels) or overt thyrotoxicosis of any etiology and in patients with acute myocardial infarction. Levothyroxine is contraindicated in patients with uncorrected adrenal insufficiency since thyroid hormones may precipitate an acute adrenal crisis by increasing the metabolic clearance of glucocorticoids (see **PRECAUTIONS**). LEVOXYL® is contraindicated in patients with hypersensitivity to any of the inactive ingredients in LEVOXYL® tablets (see **DESCRIPTION, Inactive Ingredients**).

### WARNINGS

**WARNING: Thyroid hormones, including LEVOXYL®, either alone or with other therapeutic agents, should not be used for the treatment of obesity or for weight loss. In euthyroid patients, doses within the range of daily hormonal requirements are ineffective for weight reduction. Larger doses may produce serious or even life threatening manifestations of toxicity, particularly when given in association with sympathomimetic amines such as those used for their anorectic effects.**

Levothyroxine sodium should not be used in the treatment of male or female infertility unless this condition is associated with hypothyroidism.

In patients with nontoxic diffuse goiter or nodular thyroid disease, particularly the elderly or those with underlying cardiovascular disease, levothyroxine sodium therapy is contraindicated if the serum TSH level is already suppressed due to the risk of precipitating overt thyrotoxicosis (see **CONTRAINDICATIONS**). If the serum TSH level is not suppressed, LEVOXYL® should be used with caution in conjunction with careful monitoring of thyroid function for evidence of hyperthyroidism and clinical monitoring for potential associated adverse cardiovascular signs and symptoms of hyperthyroidism.

### PRECAUTIONS

#### General

Levothyroxine has a narrow therapeutic index. Regardless of the indication for use, careful dosage titration is necessary to avoid the consequences of over- or under-treatment. These consequences include, among others, effects on growth and development, cardiovascular function, bone metabolism, reproductive function, cognitive function, emotional state, gastrointestinal function, and on glucose and lipid metabolism. Many drugs interact with levothyroxine sodium necessitating adjustments in dosing to maintain therapeutic response (see **Drug Interactions**).

**Effects on bone mineral density** – In women, long-term levothyroxine sodium therapy has been associated with decreased bone mineral density, especially in postmenopausal women on greater than replacement doses or in women who are receiving suppressive doses of levothyroxine sodium. Therefore, it is recommended that patients receiving levothyroxine sodium be given the minimum dose necessary to achieve the desired clinical and biochemical response.

**Patients with underlying cardiovascular disease** – Exercise caution when administering levothyroxine to patients with cardiovascular disorders and to the elderly in whom there is an increased risk of occult cardiac disease. In these patients, levothyroxine therapy should be initiated at lower doses than those recommended in younger individuals or in patients without cardiac disease (see **WARNINGS, PRECAUTIONS, Geriatric Use; and DOSAGE AND ADMINISTRATION**). If cardiac symptoms develop or worsen, the levothyroxine dose should be reduced or withheld for one week and then cautiously restarted at a lower dose. Overtreatment with levothyroxine sodium may have adverse cardiovascular effects such as an increase in heart rate, cardiac wall thickness, and cardiac contractility and may precipitate angina or arrhythmias. Patients with coronary artery disease who are receiving levothyroxine therapy should be monitored closely during surgical procedures, since the possibility of precipitating cardiac arrhythmias may be greater in those treated with levothyroxine. Concomitant administration of levothyroxine and sympathomimetic agents to patients with coronary artery disease may precipitate coronary insufficiency.

**Patients with nontoxic diffuse goiter or nodular thyroid disease** – Exercise caution when administering levothyroxine to patients with nontoxic diffuse goiter or nodular thyroid disease in order to prevent precipitation of thyrotoxicosis (see **WARNINGS**). If the serum TSH is already suppressed, levothyroxine sodium should not be administered (see **Contraindications**).

#### Associated endocrine disorders

**Hypothalamic/pituitary hormone deficiencies** – In patients with secondary or tertiary hypothyroidism, additional hypothalamic/pituitary hormone deficiencies should be considered, and, if diagnosed, treated (see **PRECAUTIONS, Autoimmune polyglandular syndrome**) for adrenal insufficiency.

**Autoimmune polyglandular syndrome** – Occasionally, chronic autoimmune thyroiditis may occur in association with other autoimmune disorders such as adrenal insufficiency, pernicious anemia, and insulin-dependent diabetes mellitus. Patients with concomitant adrenal insufficiency should be treated with replacement glucocorticoids prior to initiation of treatment with levothyroxine sodium. Failure to do so may precipitate an acute adrenal crisis when thyroid hormone therapy is initiated, due to increased metabolic clearance of glucocorticoids by thyroid hormone. Patients with diabetes mellitus may require upward adjustments of their antidiabetic therapeutic regimens when treated with levothyroxine (see **PRECAUTIONS, Drug Interactions**).

#### Other associated medical conditions

Infants with congenital hypothyroidism appear to be at increased risk for other congenital anomalies, with cardiovascular anomalies (pulmonary stenosis, atrial septal defect, and ventricular septal defect,) being the most common association.

#### Information for Patients

Patients should be informed of the following information to aid in the safe and effective use of LEVOXYL®:

- Notify your physician if you are allergic to any foods or medicines, are pregnant or intend to become pregnant, are breast-feeding or are taking any other medications, including prescription and over-the-counter preparations.
- Notify your physician of any other medical conditions you may have, particularly heart disease, diabetes, clotting disorders, and adrenal or pituitary gland problems. Your dose of medications used to control these other conditions may need to be adjusted while you are taking LEVOXYL®. If you have diabetes, monitor your blood and/or urinary glucose levels as directed by your physician and immediately report any changes to your physician. If you are taking anticoagulants (blood thinners), your clotting status should be checked frequently.
- Use LEVOXYL® only as prescribed by your physician. Do not discontinue or change the amount you take or how often you take it, unless directed to do so by your physician.
- The levothyroxine in LEVOXYL® is intended to replace a hormone that is normally produced by your thyroid gland. Generally, replacement therapy is to be taken for life, except in cases of transient hypothyroidism, which is usually associated with an inflammation of the thyroid gland (thyroiditis).
- Take LEVOXYL® in the morning on an empty stomach, at least one-half hour before eating any food.
- LEVOXYL® may rapidly swell and disintegrate resulting in choking, gagging, the tablet getting stuck in your throat or difficulty swallowing. It is very important that you take the tablet with a full glass of water. Most of these problems disappeared when Levoxy® tablets were taken with water.
- It may take several weeks before you notice an improvement in your symptoms.
- Notify your physician if you experience any of the following symptoms: rapid or irregular heartbeat, chest pain, shortness of breath, leg cramps, headache, nervousness, irritability, sleeplessness, tremors, change in appetite, weight gain or loss, vomiting, diarrhea, excessive sweating, heat intolerance, fever, changes in menstrual periods, hives or skin rash, or any other unusual medical event.
- Notify your physician if you become pregnant while taking LEVOXYL®. It is likely that your dose of LEVOXYL® will need to be increased while you are pregnant.
- Notify your physician or dentist that you are taking LEVOXYL® prior to any surgery.
- Partial hair loss may occur rarely during the first few months of LEVOXYL® therapy, but this is usually temporary.
- LEVOXYL® should not be used as a primary or adjunctive therapy in a weight control program.
- Keep LEVOXYL® out of the reach of children. Store LEVOXYL® away from heat, moisture, and light.

#### Laboratory Tests

##### General

The diagnosis of hypothyroidism is confirmed by measuring TSH levels using a sensitive assay (second generation assay sensitivity 0.1 mIU/L or third generation assay sensitivity ≤ 0.01 mIU/L) and measurement of free-T<sub>4</sub>.

The adequacy of therapy is determined by periodic assessment of appropriate laboratory tests and clinical evaluation. The choice of laboratory tests depends on various factors including the etiology of the underlying thyroid disease, the presence of concomitant medical conditions (including pregnancy), and the use of concomitant medications (see **PRECAUTIONS, Drug Interactions and Drug-Laboratory Test Interactions**). Persistent clinical and laboratory evidence of hypothyroidism despite an apparent adequate replacement dose of LEVOXYL® may be evidence of inadequate absorption, poor compliance, drug interactions, or decreased T<sub>4</sub> potency of the drug product.

### Adults

In adult patients with primary (thyroidal) hypothyroidism, serum TSH levels (using a sensitive assay) alone may be used to monitor therapy. The frequency of TSH monitoring during levothyroxine dose titration depends on the clinical situation but it is generally recommended at 6–8 week intervals until normalization. For patients who have recently initiated levothyroxine therapy and whose serum TSH has normalized, or in patients who have had their dosage or brand of levothyroxine changed, the serum TSH concentration should be measured after 8–12 weeks. When the optimum replacement dose has been attained, clinical (physical examination) and biochemical monitoring may be performed every 6–12 months, depending on the clinical situation, and whenever there is a change in the patient's status. It is recommended that a physical examination and a serum TSH measurement be performed at least annually in patients receiving LEVOXYL® (see **WARNINGS, PRECAUTIONS, and DOSAGE AND ADMINISTRATION**).

#### Pediatrics

In patients with congenital hypothyroidism, the adequacy of replacement therapy should be assessed by measuring both serum TSH (using a sensitive assay) and total- or free-T<sub>4</sub>. During the first three years of life, the serum total- or free-T<sub>4</sub> should be maintained at all times in the upper half of the normal range. While the aim of therapy is to also normalize the serum TSH level, this is not always possible in a small percentage of patients, particularly in the first few months of therapy. TSH may not normalize due to a resetting of the pituitary-thyroid feedback threshold as a result of *in utero* hypothyroidism. Failure of the serum T<sub>4</sub> to increase into the upper half of the normal range within 2 weeks of initiation of LEVOXYL® therapy and/or of the serum TSH to decrease below 20 mIU/L within 4 weeks should alert the physician to the possibility that the child is not receiving adequate therapy. Careful inquiry should then be made regarding compliance, dose of medication administered, and method of administration prior to raising the dose of LEVOXYL®.

The recommended frequency of monitoring of TSH and total or free T<sub>4</sub> in children is as follows: at 2 and 4 weeks after the initiation of treatment; every 1–2 months during the first year of life; every 2–3 months between 1 and 3 years of age; and every 3 to 12 months thereafter until growth is completed. More frequent intervals of monitoring may be necessary if poor compliance is suspected or abnormal values are obtained. It is recommended that TSH and T<sub>4</sub> levels, and a physical examination, if indicated, be performed 2 weeks after any change in LEVOXYL® dosage. Routine clinical examination, including assessment of mental and physical growth and development, and bone maturation, should be performed at regular intervals (see **PRECAUTIONS, Pediatric Use and DOSAGE AND ADMINISTRATION**).

#### Secondary (pituitary) and tertiary (hypothalamic) hypothyroidism

In patients with secondary or tertiary hypothyroidism, serum T<sub>4</sub> levels, which should be maintained in the upper half of the normal range in these patients.

### Drug Interactions

Many drugs affect thyroid hormone pharmacokinetics and metabolism (e.g., absorption, synthesis, secretion, catabolism, protein binding, and target tissue response) and may alter the therapeutic response to LEVOXYL®. In addition, thyroid hormones and thyroid status have varied effects on the pharmacokinetics and action of other drugs. A listing of drug-thyroidal axis interactions is contained in Table 2.

The list of drug-thyroidal axis interactions in Table 2 may not be comprehensive due to the introduction of new drugs that interact with the thyroidal axis or the discovery of previously unknown interactions. The prescriber should be aware of this fact and should consult appropriate reference sources. (e.g., package inserts of newly approved drugs, medical literature) for additional information if a drug-drug interaction with levothyroxine is suspected.

| Table 2: Drug – Thyroidal Axis Interactions   |   |   |
|---|---|---|
| Drug or Drug Class  | Effect  |   |
| <b>Drugs that may reduce TSH secretion – the reduction is not sustained; therefore, hypothyroidism does not occur</b>   |   |   |
| Dopamine / Dopamine Agonists<br>Glucocorticoids<br>Octreotide   | Use of these agents may result in a transient reduction in TSH secretion when administered at the following doses: Dopamine (> 1 mcg/kg/min); Glucocorticoids (hydrocortisone ≥ 100 mg/day or equivalent); Octreotide (> 100 mcg/day).  |   |
| <b>Drugs that alter thyroid hormone secretion</b>   |   |   |
| <b>Drugs that may decrease thyroid hormone secretion, which may result in hypothyroidism</b>  |   |   |
| Aminoglutethimide<br>Amiodarone<br>Iodide (including iodine containing Radiographic contrast agents)<br>Lithium<br>Methimazole<br>Propylthiouracil (PTU)<br>Sulfonamides<br>Tolbutamide                                       | Long-term lithium therapy can result in goiter in up to 50% of patients, and either subclinical or overt hypothyroidism, each in up to 20% of patients. The fetus, neonate, elderly and euthyroid patients with underlying thyroid disease (e.g., Hashimoto's thyroiditis or with Grave's disease previously treated with radioiodine or surgery) are among those individuals who are particularly susceptible to iodine-induced hypothyroidism. Oral cholestyralgic agents and amiodarone are slowly excreted, producing more prolonged hypothyroidism than parenterally administered iodinated contrast agents. Long-term aminoglutethimide therapy may minimally decrease T <sub>1</sub> and T <sub>4</sub> levels and increase TSH, although all values remain within normal limits in most patients. |   |
| <b>Drugs that may increase thyroid hormone secretion, which may result in hyperthyroidism</b>   |   |   |
| Amiodarone<br>Iodide (including iodine containing Radiographic contrast agents)   | Iodide and drugs that contain pharmacologic amounts of iodide may cause hyperthyroidism in euthyroid patients with Grave's disease previously treated with antithyroid drugs or in euthyroid patients with thyroid autonomy (e.g., multinodular goiter or hyperfunctioning thyroid adenoma). Hyperthyroidism may develop over several weeks and may persist for several months after therapy discontinuation. Amiodarone may induce hyperthyroidism by causing thyroiditis.   |   |
| <b>Drugs that may decrease T<sub>4</sub> absorption, which may result in hypothyroidism</b>   |   |   |
| Antacids<br>– Aluminum & Magnesium Hydroxides<br>– Simethicone<br>Bile Acid Sequestrants<br>– Cholestyramine<br>– Colestipol<br>Calcium Carbonate<br>Cation Exchange Resins<br>– Kayexalate<br>Ferrous Sulfate<br>Succralfate | Concurrent use may reduce the efficacy of levothyroxine by binding and delaying or preventing absorption, potentially resulting in hypothyroidism. Calcium carbonate may form an insoluble chelate with levothyroxine, and ferrous sulfate likely forms a ferric-thyroxine complex. Administer levothyroxine at least 4 hours apart from these agents.  |   |
| <b>Drugs that may alter T<sub>4</sub> and T<sub>3</sub> serum transport – but FT<sub>4</sub> concentration remains normal; and, therefore, the patient remains euthyroid</b>  |   |   |
| <b>Drugs that may increase serum TBG concentration</b>  |   | <b>Drugs that may decrease serum TBG concentration</b>  |
| Clofibrate<br>Estrogen-containing oral contraceptives<br>Estrogens (oral)<br>Heron / Methadone<br>5-Fluorouracil<br>Mitotane<br>Tamoxifen   | Androgens / Anabolic Steroids<br>Asparaginase<br>Glucocorticoids<br>Slow-Release Nicotinic Acid   |   |
| <b>Drugs that may cause protein-binding site displacement</b>   |   | Administration of these agents with levothyroxine results in an initial transient increase in FT <sub>4</sub> . Continued administration results in a decrease in serum T <sub>4</sub> and normal FT <sub>4</sub> and TSH concentrations and, therefore, patients are clinically euthyroid. Salicylates inhibit binding of T <sub>3</sub> and T <sub>4</sub> to TBG and thyroxin. An initial increase in serum FT <sub>4</sub> is followed by return of FT <sub>4</sub> to normal levels with sustained therapeutic serum salicylate concentrations, although total-T <sub>4</sub> levels may decrease by as much as 30%. |

| Drugs that may alter $T_4$ and $T_3$ metabolism   |  |
|---|--|
| <b>Drugs that may increase hepatic metabolism, which may result in hypothyroidism</b>   |  |
| Carbamazepine<br>Phenytoin<br>Hydrochlorothiazide<br>Rifampin   | Stimulation of hepatic microsomal drug-metabolizing enzyme activity may cause increased hepatic degradation of levvothyroxine, resulting in increased levvothyroxine requirements. Phenytoin and carbamazepine reduce serum protein binding of levvothyroxine, and total- and free- $T_4$ may be reduced by 20% to 40%, but most patients have normal serum TSH levels and are clinically euthyroid.   |
| <b>Drugs that may decrease <math>T_4</math> 5'-deiodinase activity</b>  |  |
| Amiodarone<br>Beta-adrenergic antagonists<br>- (e.g., Propranolol) >160 mg/day<br>Glucocorticoids<br>- (e.g., Dexamethasone >4 mg/day)<br>Propylthiouracil (PTU)  | Administration of these enzyme inhibitors decreases the peripheral conversion of $T_4$ to $T_3$ leading to decreased $T_3$ levels. However, serum $T_4$ levels are usually normal but may occasionally be slightly increased. In patients treated with large doses of propranolol (>160 mg/day), $T_3$ and $T_4$ levels change slightly. TSH levels are small, normal, and patients are clinically euthyroid. It should be noted that actions of particular beta-adrenergic antagonists may be impaired when the hypothyroid patient is converted to the euthyroid state. Short-term administration of large doses of glucocorticoids may decrease serum $T_3$ concentrations by 30% with minimal change in serum $T_4$ levels. However, long-term glucocorticoid therapy may result in slightly decreased $T_3$ and $T_4$ levels due to decreased TBG production (see above). |
| <b>Miscellaneous</b>  |  |
| Anticoagulants (oral)<br>- Coumatin Derivatives<br>- Indandione Derivatives   | Thyroid hormones appear to increase the catabolism of vitamin K-dependent clotting factors, thereby increasing the anticoagulant activity of oral anticoagulants. Concomitant use of these agents impairs the compensatory increases in clotting factor synthesis. Prothrombin time should be carefully monitored in patients taking levvothyroxine and oral anticoagulants and the dose of anticoagulant therapy adjusted accordingly.  |
| Antidepressants<br>- Tricyclics (e.g., Amitriptyline)<br>- Tetracyclics (e.g., Maprotiline)<br>- Selective Serotonin Reuptake Inhibitor (SSRIs; e.g., Sertraline)   | Concurrent use of tricyclic antidepressants and levvothyroxine may increase the therapeutic and toxic effects of both drugs, possibly due to increased receptor sensitivity to catecholamines. Toxic effects may include increased risk of cardiac arrhythmias and CNS stimulation; onset of action of tricyclics may be accelerated. Administration of serotonin reuptake inhibitors on levvothyroxine may result in increased levvothyroxine requirements.   |
| Antidiabetic Agents<br>- Biguanides<br>- Meglitinides<br>- Sulfonylureas<br>- Thiazolidinediones<br>- Insulin   | Addition of levvothyroxine to antidiabetic or insulin therapy may result in increased antidiabetic agent or insulin requirements. Careful monitoring of diabetic control is recommended, especially when thyroid therapy is started, changed, or discontinued.   |
| Cardiac Glycosides  | Serum digitalis glycoside levels may be reduced in hypothyroidism or when the hypothyroid patient is converted to the euthyroid state. Therapeutic effect of digitalis glycosides may be reduced.  |
| Cytokines<br>- Interferon- $\alpha$<br>- Interleukin-2  | Therapy with interferon- $\alpha$ has been associated with the development of antithyroid microsomal antibodies in 20% of patients and some have transient hypothyroidism, hyperthyroidism, or both. Patients who have antithyroid antibodies before treatment are at higher risk for thyroid dysfunction during treatment. Interleukin-2 has been associated with transient painless thyroiditis in 20% of patients. Interferon- $\beta$ and - $\gamma$ have not been reported to cause thyroid dysfunction.  |
| Growth Hormones<br>- Somatrem<br>- Somatropin   | Excessive use of thyroid hormones with growth hormones may accelerate epiphyseal closure. However, untreated hypothyroidism may interfere with growth response to growth hormone.  |
| Ketamine  | Concurrent use may produce marked hypertension and tachycardia; cautious administration to patients receiving thyroid hormone therapy is recommended.  |
| Methylxanthine Bronchodilators<br>- (e.g., Theophylline)  | Decreased theophylline clearance may occur in hypothyroid patients; clearance returns to normal when the euthyroid state is achieved.  |
| Radiographic Agents   | Thyroid hormones may reduce the uptake of $^{131}\text{I}$ , $^{125}\text{I}$ , and $^{99\text{m}}\text{Tc}$ .   |
| Sympathomimetics  | Concurrent use may increase the effects of sympathomimetics on thyroid hormone. Thyroid hormones may increase the risk of coronary insufficiency when sympathomimetic agents are administered to patients with coronary artery disease.  |
| Chloral Hydrate<br>Diazepam<br>Ethionamide<br>Lovastatin<br>Metoclopramide<br>6-Mercaptopurine<br>Nitroglycerine<br>Para-aminosalicylate sodium<br>Perphenazine<br>Resorcinol (excessive topical use)<br>Thiazide Diuretics | These agents have been associated with thyroid hormone and/or TSH level alterations by various mechanisms.   |

**Oral anticoagulants** – Levvothyroxine increases the response to oral anticoagulant therapy. Therefore, a decrease in the dose of anticoagulant may be warranted with correction of the hypothyroid state or when the LEVOXYL<sup>®</sup> dose is increased. Prothrombin time should be closely monitored to permit appropriate and timely dosage adjustments (see Table 2).

**Digitalis glycosides** – The therapeutic effects of digitalis glycosides may be reduced by levvothyroxine. Serum digitalis glycoside levels may be decreased when a hypothyroid patient becomes euthyroid, necessitating an increase in the dose of digitalis glycosides (see Table 2).

**Drug-Food Interactions** – Consumption of certain foods may affect levvothyroxine absorption thereby necessitating adjustments in dosing. Soybean flour, cornmeal, cotton seeds, walnuts, and dietary fiber may bind and decrease the absorption of levvothyroxine sodium from the GI tract.

**Drug-Laboratory Test Interactions** – Changes in TBG concentration must be considered when interpreting  $T_4$  and  $T_3$  values, which necessitates measurement and evaluation of unbound (free) hormone and/or determination of the free  $T_4$  index (FT $_4$ ). Pregnancy, infectious hepatitis, estrogens, estrogen-containing oral contraceptives, and acute intermittent porphyria increase TBG concentrations. Decreases in TBG concentrations are observed in nephrosis, severe hypoproteinemia, severe liver disease, acromegaly, and after androgen or corticosteroid therapy (see also Table 2). Familial hyper- or hypothyroxinemia binding globulinemias have been described, with the incidence of TBG deficiency approximating 1 in 5000.

**Carcinogenesis, Mutagenesis, and Impairment of Fertility** – Animal studies have not been performed to evaluate the carcinogenic potential, mutagenic potential or effects on fertility of levvothyroxine. The synthetic  $T_4$  in LEVOXYL<sup>®</sup> is identical to that produced naturally by the human thyroid gland. Although there has been a reported association between prolonged thyroid hormone therapy and breast cancer, this has not been confirmed. Patients receiving LEVOXYL<sup>®</sup> for appropriate clinical indications should be treated to the lowest effective replacement dose.

**Pregnancy – Category A** – Studies in women taking levvothyroxine sodium during pregnancy have not shown an increased risk of congenital abnormalities. Therefore, the possibility of fetal harm appears remote. LEVOXYL<sup>®</sup> should not be discontinued during pregnancy and hypothyroidism diagnosed during pregnancy should be promptly treated.

Hypothyroidism during pregnancy is associated with a higher rate of complications, including spontaneous abortion, pre-eclampsia, stillbirth and premature delivery. Maternal hypothyroidism may have an adverse effect on fetal and childhood growth and development. During pregnancy, serum  $T_4$  levels may decrease and serum TSH levels increase to values outside the normal range. Since elevations in serum TSH may occur as early as 4 weeks gestation, pregnant women taking LEVOXYL<sup>®</sup> should have their TSH measured during each trimester. An elevated serum TSH level should be corrected by an increase in the LEVOXYL<sup>®</sup> dose. Postpartum TSH levels are similar to preconception values, the LEVOXYL<sup>®</sup> dose should return to the pre-pregnancy dose immediately after delivery. A serum TSH level should be obtained 6–8 weeks postpartum.

Thyroid hormones do not readily cross the placental barrier; however, some transfer does occur as evidenced by levels in a cord blood of athletic fetuses being approximately one-third maternal levels. Transfer of thyroid hormone from the mother to the fetus, however, may not be adequate to prevent in utero hypothyroidism.

**Nursing Mothers** – Although thyroid hormones are excreted only minimally in human milk, caution should be exercised when LEVOXYL<sup>®</sup> is administered to a nursing woman. However, adequate replacement doses of levvothyroxine are generally needed to maintain normal lactation.

#### Pediatric Use

##### General

The goal of treatment in pediatric patients with hypothyroidism is to achieve and maintain normal intellectual and physical growth and development.

The initial dose of levvothyroxine varies with age and body weight (see DOSE AND ADMINISTRATION, Table 3). Dosing adjustments are based on an assessment of the individual patient's clinical and laboratory parameters (see PRECAUTIONS, Laboratory Tests).

In children in whom a diagnosis of permanent hypothyroidism has not been established, it is recommended that levvothyroxine administration be discontinued for a 30-day trial period, but only after the child is at least 3 years of age. Serum  $T_4$  and TSH levels should then be obtained. If the  $T_4$  is low and the TSH high, the diagnosis of permanent hypothyroidism is established, and levvothyroxine therapy should be reinitiated. If the  $T_4$  and TSH levels are normal, euthyroidism may be assumed and, therefore, the hypothyroidism is considered to have been transient. In this instance, however, the physician should carefully monitor the child and repeat the thyroid function tests if any signs or symptoms of hypothyroidism develop. In this setting, the clinician should have a high index of suspicion of relapse. If the results of the levvothyroxine withdrawal test are inconclusive, careful follow-up and subsequent testing will be necessary.

Since more severely affected children may become clinically hypothyroid when treatment is discontinued for 30 days, an alternate approach is to reduce the replacement dose of levvothyroxine by half during the 30-day trial period. If, after 30 days, the serum TSH is elevated above 20 mIU/L, the diagnosis of permanent hypothyroidism is confirmed, and full replacement therapy should be resumed. However, if the serum TSH has not risen to greater than 20 mIU/L, levvothyroxine treatment should be discontinued for another 30-day trial period followed by repeat serum  $T_4$  and TSH.

The presence of concomitant medical conditions should be considered in certain clinical circumstances and, if present, appropriate treatment should be initiated (see PRECAUTIONS, Congenital Hypothyroidism [see PRECAUTIONS, Laboratory Tests and DOSE AND ADMINISTRATION]). Rapid restoration of normal serum  $T_4$  concentrations is essential for preventing the adverse effects of congenital hypothyroidism on intellectual development as well as on overall physical growth and maturation. Therefore, LEVOXYL<sup>®</sup> therapy should be initiated immediately upon diagnosis and is generally continued for life.

During the first 2 weeks of LEVOXYL<sup>®</sup> therapy, infants should be closely monitored for cardiac overload, arrhythmias, and aspiration from adult suckling.

The patient should be monitored closely to avoid undertreatment or overtreatment. Undertreatment may have deleterious effects on intellectual development and skeletal growth. Overtreatment has been associated with craniosynostosis in infants, and may adversely affect the tempo of brain maturation and accelerate the bone age with resultant premature closure of the epiphyses and compromised adult stature.

**Acquired Hypothyroidism in Pediatric Patients.**

The patient should be monitored closely to avoid undertreatment and overtreatment. Undertreatment may result in poor school performance due to impaired concentration and slowed mentation and in reduced adult height. Overtreatment may accelerate the bone age and result in premature epiphyseal closure and compromised adult stature.

Treated children may manifest a period of catch-up growth, which may be adequate in some cases to normalize adult height. In children with severe or prolonged hypothyroidism, catch-up growth may not be adequate to normalize adult height.

#### Geriatric Use

Because of the increased prevalence of cardiovascular disease among the elderly, levvothyroxine therapy should not be initiated at the full replacement dose (see WARNINGS, PRECAUTIONS, and DOSE AND ADMINISTRATION).

#### ADVERSE REACTIONS

Adverse reactions associated with levvothyroxine therapy are primarily those of hyperthyroidism due to therapeutic overdose. They include the following:

**General:** fatigue, increased appetite, weight loss, heat intolerance, fever, excessive sweating.

**Central nervous system:** headache, hyperactivity, nervousness, anxiety, irritability, emotional lability, insomnia.

**Musculoskeletal:** tremors, muscle weakness.

**Cardiac:** palpitations, tachycardia, arrhythmias, increased pulse and blood pressure, heart failure, angina, myocardial infarction, cardiac arrest.

**Pulmonary:** dyspnea.

**GI:** diarrhea, vomiting, abdominal cramps.

**Dermatologic:** hair loss, flushing.

**Reproductive:** menstrual irregularities, impaired fertility.

Pseudotumor cerebri and slipped capital femoral epiphysis have been reported in children receiving levvothyroxine therapy. Overtreatment may result in craniosynostosis in infants and premature closure of the epiphyses in children with resultant compromised adult height.

Seizures have been reported rarely with the institution of levvothyroxine therapy. Inadequate levvothyroxine dosage will produce or fail to ameliorate the signs and symptoms of hypothyroidism.

Hypersensitivity reactions to inactive ingredients have occurred in patients treated with thyroid hormone products. These include urticaria, pruritus, skin rash, flushing, angioedema, various GI symptoms (abdominal pain, nausea, vomiting and diarrhea), fever, arthralgia, serum sickness and wheezing. Hypersensitivity to thyroid hormone is not known to occur.

In addition to the above events, the following have been reported, presumably when Levoxy<sup>®</sup> tablets were not taken with water: choking, gagging, tablet stuck in throat and dysphagia (see Information for Patients).

#### OVERDOSAGE

The signs and symptoms of overdose are those of hyperthyroidism (see PRECAUTIONS and ADVERSE REACTIONS). In addition, confusion and disorientation may occur. Cerebral embolism, shock, coma, and death have been reported. Seizures have occurred in a child ingesting approximately 20 mg of levvothyroxine. Symptoms may not necessarily be evident or may not appear until several days after ingestion of levvothyroxine sodium.

**Treatment of Overdosage**

Levvothyroxine overdose should be reduced in dose or temporarily discontinued if signs or symptoms of overdose occur.

**Acute Massive Overdosage** – This may be a life-threatening emergency, therefore, symptomatic and supportive therapy should be instituted immediately. If not contraindicated (e.g., by seizures, coma, or loss of the gag reflex), the stomach should be emptied by emesis or gastric lavage to decrease gastrointestinal absorption. Activated charcoal or cholestyramine may also be used to decrease absorption. Central and peripheral increased sympathetic activity may be treated by administering  $\beta$ -receptor antagonists, e.g., propranolol (1 to 3 mg intravenously over a 10-minute period, or orally, 80 to 160 mg/day). Provide respiratory support as needed; control congestive heart failure; control fever; hypokalemia, and fluid loss as necessary. Glucocorticoids may be given to inhibit the conversion of  $T_4$  to  $T_3$ . Because  $T_4$  is highly protein bound, very little drug will be removed by dialysis.

#### DOSE AND ADMINISTRATION

##### General Principles:

The goal of replacement therapy is to achieve and maintain a clinical and biochemical euthyroid state. The goal of suppressive therapy is to inhibit growth and/or function of abnormal thyroid tissue. The dose of LEVOXYL<sup>®</sup> that is adequate to achieve these goals depends on a variety of factors including the patient's age, body weight, cardiovascular status, concomitant medical conditions, including pregnancy, concomitant medications, and the specific nature of the condition being treated (see WARNINGS and PRECAUTIONS). Hence, the following recommendations serve only as dosing guidelines. Dosing must be individualized and adjustments made based on periodic assessment of the patient's clinical response and laboratory parameters (see PRECAUTIONS, Laboratory Tests).

The LEVOXYL<sup>®</sup> should be taken in the morning on an empty stomach, at least one-half hour before any food is eaten. LEVOXYL<sup>®</sup> should be taken at least 4 hours apart from drugs that are known to interfere with its absorption (see PRECAUTIONS, Drug Interactions).

LEVOXYL<sup>®</sup> should be taken with water (see Information for Patients and ADVERSE REACTIONS).

Due to the long half-life of levvothyroxine, the peak therapeutic effect at a given dose of levvothyroxine sodium may not be attained for 4–6 weeks.

Caution should be exercised when administering LEVOXYL<sup>®</sup> to patients with underlying cardiovascular disease, to the elderly, and to those with concomitant adrenal insufficiency (see PRECAUTIONS, Specific Patient Populations).

**Hypothyroidism in Adults and in Children in Whom Growth and Puberty are Complete (see WARNINGS and PRECAUTIONS, Laboratory Tests)**

Thyroid therapy should be initiated in otherwise healthy individuals less than 50 years old and in those older than 50 years who have been recently treated for hypothyroidism or who have been hypothyroid for only a short time (such as a few months). The average full replacement dose of levvothyroxine sodium is approximately 1.7 mcg/kg/day (e.g., 100–125 mcg/day for a 70 kg adult). Older patients may require less than 1 mcg/kg/day. Levvothyroxine sodium doses greater than 200 mcg/day are seldom required. An inadequate response to daily doses 300 mcg/day is rare and may indicate poor compliance, malabsorption, and/or drug interactions.

For most patients older than 50 years or for patients under 50 years of age with underlying cardiac disease, an initial starting dose of 25–50 mcg/day of levvothyroxine sodium is recommended, with gradual increments in dose at 6–8 week intervals, as needed. The recommended starting dose of levvothyroxine sodium in elderly patients with cardiac disease is 12.5–25 mcg/day, with gradual dose increments at 4–6 week intervals. The levvothyroxine sodium dose is generally adjusted in 12.5–25 mcg increments until the patient with primary hypothyroidism is clinically euthyroid and the serum TSH has normalized.

In patients with severe hypothyroidism, the recommended initial levvothyroxine sodium dose is 12.5–25 mcg/day with increases of 25 mcg/day every 2–4 weeks, accompanied by clinical and laboratory assessment, until the TSH level is normalized.

In patients with secondary (pituitary) or tertiary (hypothalamic) hypothyroidism, the levvothyroxine sodium dose should be titrated until the patient is clinically euthyroid and the serum free- $T_4$  level is restored to the upper half of the normal range.

**Pediatric Use – Congenital or Acquired Hypothyroidism (see PRECAUTIONS, Laboratory Tests) General Principles**

In general, levvothyroxine therapy should be instituted at full replacement doses as soon as possible. Delays in diagnosis and institution of therapy may have deleterious effects on the child's intellectual and physical growth and development.

Undertreatment and overtreatment should be avoided (see PRECAUTIONS, Pediatric Use).

LEVOXYL<sup>®</sup> may be administered to infants and children who cannot swallow intact tablets by crushing

the tablet and suspending the freshly crushed tablet in a small amount (5–10 mL or 1–2 teaspoons) of water. This suspension can be administered by spoon or dropper. **DO NOT STORE THE SUSPENSION.** Foods that decrease absorption of levvothyroxine, such as soybean infant formula, should not be used for administering levvothyroxine sodium tablets. (see PRECAUTIONS, Drug-Food Interactions).

#### Newborns

The recommended starting dose of levvothyroxine sodium in newborn infants is 10–15 mcg/kg/day. A lower starting dose (e.g., 25 mcg/day) should be considered in infants at risk for cardiac failure, and the dose should be increased in 4–6 weeks as needed based on clinical and laboratory response to treatment. In infants with very low (<5 mcg/dL) or undetectable serum  $T_4$  concentrations, the recommended initial starting dose is 50 mcg/day of levvothyroxine sodium.

#### Infants and Children

Levvothyroxine therapy is usually initiated at full replacement doses, with the recommended dose per body weight decreasing with age (see Table 3). However, in children with chronic or severe hypothyroidism, an initial dose of 25 mcg/day of levvothyroxine sodium is recommended with increments of 25 mcg every 2–4 weeks until the desired effect is achieved.

Hyperactivity in an older child can be minimized if the starting dose is one-fourth of the recommended full replacement dose, and the dose is then increased on a weekly basis by an amount equal to one-fourth the full recommended replacement dose until the full recommended replacement dose is reached.

| AGE                         | Daily Dose Per Kg Body Weight <sup>a</sup> |
|-----------------------------|--|
| 0–3 months                  | 10–15 mcg/kg/day                           |
| 3–6 months                  | 8–10 mcg/kg/day                            |
| 6–12 months                 | 6–8 mcg/kg/day                             |
| 1–5 years                   | 5–6 mcg/kg/day                             |
| 6–12 years                  | 4–6 mcg/kg/day                             |
| >12 years                   | 2–3 mcg/kg/day                             |
| Growth and puberty complete | 1.7 mcg/kg/day                             |

<sup>a</sup> - The dose should be adjusted based on clinical response and laboratory parameters (see PRECAUTIONS, Laboratory Tests and Pediatric Use).

**Pregnancy** – Pregnancy may increase levvothyroxine requirements (see PREGNANCY).

**Subclinical Hypothyroidism** – If this condition is treated, a lower levvothyroxine sodium dose (e.g., 1 mcg/kg/day) than that used for full replacement may be adequate to normalize the serum TSH level. Patients who are not treated should be monitored yearly for changes in clinical status and thyroid laboratory parameters.

**TSH Suppression in Well-Differentiated Thyroid Cancer and Thyroid Nodules** – The target level for TSH suppression in these conditions has not been established with controlled studies. In addition, the efficacy of TSH suppression for benign nodular disease is controversial. Therefore, the dose of LEVOXYL<sup>®</sup> used for TSH suppression should be individualized based on the specific disease and the patient being treated.

In the treatment of well-differentiated (papillary and follicular) thyroid cancer, levvothyroxine is used as an adjunct to surgical and radioactive therapy. Generally, TSH is suppressed to <0.1 mIU/L and this usually requires a levvothyroxine sodium dose of greater than 2 mcg/kg/day. However, in patients with high-risk tumors, the target level for TSH suppression may be <0.01 mIU/L.

In the treatment of benign nodules and nontoxic multinodular goiter, TSH is generally suppressed to a higher target level, 0.1–0.5 mIU/L for nodules and 0.5–1.0 mIU/L for multinodular goiter) than that used for the treatment of thyroid cancer. Levvothyroxine sodium is contraindicated if the serum TSH is already suppressed due to the risk of precipitating overt hypothyroidism (see CONTRAINDICATIONS, WARNINGS and PRECAUTIONS).

**Myxedema Coma** – Myxedema coma is a life-threatening emergency characterized by poor circulation and hypometabolism, and may result in unpredictable absorption of levvothyroxine sodium from the gastrointestinal tract. Therefore, oral thyroid hormone drug products are not recommended to treat this condition. Thyroid hormone products formulated for intravenous administration should be administered.

#### HOW SUPPLIED

| Strength (mcg) | Color     | NDC # for bottles of 100 | NDC # for bottles of 1000 | NDC # for Unit Dose Cartons of 100 |
|----------------|-----------|--------------------------|---------------------------|------------------------------------|
| 25             | Orange    | NDC 52604-5025-1         | NDC 52604-5025-2          | NDC 52604-5025-5                   |
| 50             | White     | NDC 52604-5050-1         | NDC 52604-5050-2          | NDC 52604-5050-5                   |
| 75             | Purple    | NDC 52604-5075-1         | NDC 52604-5075-2          | NDC 52604-5075-5                   |
| 88             | Olive     | NDC 52604-5088-1         | NDC 52604-5088-2          | NDC 52604-5088-5                   |
| 100            | Yellow    | NDC 52604-5100-1         | NDC 52604-5100-2          | NDC 52604-5100-5                   |
| 112            | Rose      | NDC 52604-5112-1         | NDC 52604-5112-2          | NDC 52604-5112-5                   |
| 125            | Brown     | NDC 52604-5125-1         | NDC 52604-5125-2          | NDC 52604-5125-5                   |
| 137            | Dark Blue | NDC 52604-5137-1         | NDC 52604-5137-2          | NDC 52604-5137-5                   |
| 150            | Blue      | NDC 52604-5150-1         | NDC 52604-5150-2          | NDC 52604-5150-5                   |
| 175            | Turquoise | NDC 52604-5175-1         | NDC 52604-5175-2          | NDC 52604-5175-5                   |
| 200            | Pink      | NDC 52604-5200-1         | NDC 52604-5200-2          | NDC 52604-5200-5                   |
| 300            | Green     | NDC 52604-5300-1         | NDC 52604-5300-2          | NDC 52604-5300-5                   |

— LEVOXYL<sup>®</sup> (levvothyroxine sodium tablets, USP) are supplied as oval, color-coded, potency marked tablets in 12 strengths:

#### STORAGE CONDITIONS

20°–25°C (68°–77°F) with excursions permitted between 15°–30°C (59°–86°F).

Meets USP Dissolution Tests 1 and 2.

#### ML NLY

#### MANUFACTURER

JONES PHARMA INCORPORATED

(A wholly owned subsidiary of King Pharmaceuticals, Inc.)

Bristol, VA 24201

Prescribing Information as of May 2004



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Contact King Pharmaceuticals at 501 Fifth Street, Bristol, Tennessee:  
888-358-6438; or www.levoxy.com.