Lithium Associated Hyperparathyroidism: An Evidence Based Surgical Approach

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Received May 17, 2011; revised August 15, 2011; accepted October 17, 2011

Abstract

Background: Long-term lithium use in psychiatric patients may lead to lithium associated hyperparathyroidism (LAH). Although anecdotal case reports have appeared, an evidence based algorithm for management of LAH is lacking. Methods: A comprehensive literature search was performed (1973-2010) using PubMed with keywords; “lithium” “hypercalcemia” “hyperparathyroidism” “sestamibi” “intra-operative parathyroid hormone (IOPTH) monitoring” “parathyroidectomy” and “medical management”. All English language publications addressing etiology and clinical management issues concerning LAH were critically analyzed. Results: Lithium associated hyperparathyroidism occurs in 4.3% - 6.3% of chronic lithium users compared to the general population which has an incidence of 0.5% - 1%. 194 cases of LAH have been reported which includes 10 patients (5%) treated medically and 170 patients (88%) who underwent parathyroidectomy. Details were available for 14 patients (7%). Among parathyroidectomy patients, 104 (59%) had adenomatous disease and 66 (39%) had multiglandular hyperplasia. Preoperative localization studies were utilized in only 22 patients (13%) and IOPTH monitoring was reported in only 3 studies (32 patients, 19%). Among surgical patients, bilateral neck exploration (BNE) was the most common approach performed in 162 patients (95%); focused neck exploration was utilized in only 8 patients (5%). Parathyroidectomy normalized LAH biochemical changes in nearly all patients (90% - 97%) in the early post-operative period, but recurrent hyperparathyroidism occurred in 8% - 42% of patients. Conclusion: LAH is an under appreciated and poorly understood endocrine disorder. LAH has a higher incidence of multiglandular disease and bilateral neck exploration is mandatory in majority for disease control. Nonsurgical approaches may be useful in select patients on short-term lithium therapy.

Keywords: Lithium, Hypercalcemia, Hyperparathyroidism, Sestamibi Intra-Operative Parathyroid Hormone (IOPTH) Monitoring, Parathyroidectomy, Medical Management

1. Introduction

Lithium Carbonate (Li⁺) is the preferred and most efficacious therapy for acute treatment and maintenance therapy for bipolar depressive disorder, and is a useful adjunct in unipolar depression [1,2]. Long-term lithium therapy is associated with multiple endocrine and metabolic alterations such as hyperthyroidism, hypothyroidism, reduced bone mineral density (BMD), osteopenia, as well as several gastrointestinal (nausea, constipation), cardiovascular (bradyarrhythmias), renal (nephrocalcinosis, reduced glomerular filtration rate (GFR), polyuria, nephrogenic diabetes insipidus) and psychosomatic adverse effects (weakness, fatigue, depression). Chronic lithium therapy is also associated with hyperparathyroidism (LAH) characterized by hypercalcemia, hypermagnesemia, reduced urinary calcium, and elevated serum PTH levels often with enlarged parathyroid gland(s) [3,4]. Rare case reports and a limited number of reviews have yielded inadequate and conflicting data concerning the ideal treatment of LAH [4-6]. This article provides a systematic review of LAH, addresses current controversies, and pro-
poses an algorithmic approach to LAH treatment.

1.1. Methods/Materials

All English language LAH publications for the years 1973 to 2010 which addressed the epidemiology, biochemical association, etiology, pathogenesis, clinical presentation, preoperative imaging studies, intraoperative parathyroid hormone monitoring (IOPTH), surgical management (parathyroidectomy) and surveillance were collected and analyzed. The literature search was performed utilizing PubMed with the following keywords; “lithium” “hypercalcemia” “hyperparathyroidism” “sestamibi” “IOPTH” “parathyroidectomy” “medical management” and “cinacalcet”.

1.2. Results

Lithium use is associated with an increased incidence of hyperparathyroidism (4.3% - 6.3%) and a female preponderance (F: M 4:1) compared to primary hyperparathyroidism in the general population (0.5% - 1% and F: M 3:1). A total of 194 cases of LAH have been reported in English language literature of which 170 patients (88%) were treated by surgical resection (parathyroidectomy), 10 patients (5%) were managed medically and no details are available for 14 patients (7%). (Tables 1 and 2) Among those who had parathyroidectomy, 104 patients (61%) had adenomatous disease and 66 patients (39%) had multigland parathyroid hyperplasia compared to primary hyperparathyroidism (PHPT) in the general population in whom single adenoma is seen in 85% and multigland hyperplasia is identified in 15%. Among 170 LAH patients undergoing parathyroidectomy, preoperative localization studies (sestamibi scan, neck ultrasound, CT scan) were utilized in only 22 patients (14%) and IOPTH monitoring was reported in only 3 studies (32 patients, 19%).

Bilateral neck exploration was the most common surgical approach utilized and was utilized in 162 patients (95%), whereas focused neck exploration was performed

<table>
<thead>
<tr>
<th>Author, Year</th>
<th>Total (N =)</th>
<th>Surgically Treated (N =) (%)</th>
<th>Adenoma</th>
<th>Four gland hyperplasia (N =) (%)</th>
<th>Remarks (Surgical approach, BNE, IOPTH monitoring)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Garfinkel et al. [10], 1973</td>
<td>1</td>
<td>1 (100%)</td>
<td>1 (100%)</td>
<td>-</td>
<td>First reported case, BNE</td>
</tr>
<tr>
<td>Ananth et al. [20], 1983</td>
<td>3</td>
<td>2 (67%)</td>
<td>2 (67%)</td>
<td>-</td>
<td>BNE, 1 patient-details-NA</td>
</tr>
<tr>
<td>Stancer et al. [15], 1989</td>
<td>8</td>
<td>3 (37.5%)</td>
<td>1 (12.5%)</td>
<td>2 (25%)</td>
<td>BNE, 5 patients-details-NA</td>
</tr>
<tr>
<td>Nordenstrom et al. [19], 1992</td>
<td>6</td>
<td>6 (100%)</td>
<td>1 (17%)</td>
<td>5 (83%)</td>
<td>BNE</td>
</tr>
<tr>
<td>Bendz et al. [8], 1996</td>
<td>8</td>
<td>8 (100%)</td>
<td>3 (37.5%)</td>
<td>2 (25%)</td>
<td>Point prevalence of LAH-3.6%</td>
</tr>
<tr>
<td>McHenry et al. [3], 1996</td>
<td>25</td>
<td>17 (68%)</td>
<td>12 (48%)</td>
<td>5 (20%)</td>
<td>BNE, 8 patients-details-NA</td>
</tr>
<tr>
<td>Wolf et al. [21], 1997</td>
<td>1</td>
<td>1 (100%)</td>
<td>-</td>
<td>1 (100%)</td>
<td>BNE</td>
</tr>
<tr>
<td>deCelis et al. [22], 1998</td>
<td>1</td>
<td>1 (100%)</td>
<td>1 (100%)</td>
<td>-</td>
<td>BNE</td>
</tr>
<tr>
<td>Abdullah et al. [17], 1999</td>
<td>11</td>
<td>11 (100%)</td>
<td>6 (55%)</td>
<td>3 (27%)</td>
<td>BNE, 1 recurrence</td>
</tr>
<tr>
<td>Awad et al. [5], 1999</td>
<td>15</td>
<td>15 (100%)</td>
<td>11 (73%)</td>
<td>3 (20%)</td>
<td>BNE</td>
</tr>
<tr>
<td>Hundley et al. [18], 2005</td>
<td>12</td>
<td>12 (100%)</td>
<td>6 (50%)</td>
<td>3 (25%)</td>
<td>IOPTH monitoring</td>
</tr>
<tr>
<td>Carchman et al. [16], 2008</td>
<td>16</td>
<td>16 (100%)</td>
<td>12 (75%)</td>
<td>4 (25%)</td>
<td>IOPTH monitoring, 8 patients had focused neck exploration</td>
</tr>
<tr>
<td>Rizwan et al. [27], 2009</td>
<td>1</td>
<td>1 (100%)</td>
<td>-</td>
<td>1 (100%)</td>
<td>BNE</td>
</tr>
<tr>
<td>Szalat et al. [12], 2008</td>
<td>4</td>
<td>4 (100%)</td>
<td>1 (25%)</td>
<td>2 (50%)</td>
<td>75% MGH, IOPTH monitoring</td>
</tr>
<tr>
<td>Jarhult et al. [14], 2010</td>
<td>71</td>
<td>71 (100%)</td>
<td>32 (45%)</td>
<td>37 (52%)</td>
<td>BNE, 42% had recurrence</td>
</tr>
<tr>
<td>Current study, 2010</td>
<td>1</td>
<td>1 (100%)</td>
<td>-</td>
<td>1 (100%)</td>
<td>IOPTH, BNE</td>
</tr>
<tr>
<td>Overall</td>
<td>184*</td>
<td>170 (92%)</td>
<td>89 (52%)</td>
<td>66 (39%)</td>
<td></td>
</tr>
</tbody>
</table>

Abbreviations: N, number of patients; BNE, bilateral neck exploration; MGH, Multiglandular hyperplasia; IOPTH, intraoperative parathyroid hormone monitoring; LAH, lithium associated hyperparathyroidism; NA, not available. Note: Published studies on the surgical management of LAH document that a single adenoma is the most common LAH associated pathology (61%). However, increased prevalence of double adenomas (7%) and multigland hyperplasia (39%) suggests that bilateral neck exploration is necessary in the majority of LAH patients (96%). Pre-operative localization studies and IOPTH monitoring may have utility in LAH patients, however data on their value is limited (N = 22, 33 respectively).
in 8 (5%) patients. The latter approach was utilized significantly less often in LAH when compared to PHPT patients in whom the published unilateral focused neck exploration rate is 80% - 95% [7]. This change likely reflects the increased incidence of four-gland hyperplasia in LAH. Parathyroidectomy normalized LAH associated biochemical changes in nearly all patients (67% - 100%) in the early post-operative period, however at a median long-term follow-up of 6.3 years 8% - 42% patients experienced recurrence.

### 2. Case Review

A 64 year-old female with a past medical history of type II diabetes mellitus, hypertension, hypercholesterolemia, and bipolar depression was referred for surgical evaluation due to persistent hypercalcemia. She had been treated with lithium (300 mg BID) for four years. Over the prior three years, the patient had mild elevations of her serum calcium level (normal range; 10 - 11 mg/dL) which rose to 11.7 mg/dL and 12.3 mg/dL over a two months duration. Symptomatically she complained of decreased memory and mental clarity. Her diabetes was well controlled (hemoglobin A1C level of 6%) with no evidence of retinopathy or nephropathy. A sestamibi parathyroid scan localized a parathyroid adenoma at the mid pole of the left lobe of the thyroid gland. The patient was scheduled for a PTH directed left parathyroidectomy under local anesthesia. Pre-operative PTH level was 147 pg/ml (normal range; 10 - 55 pg/ml). At exploration the large cystic structure was identified on the anterior surface of the left lower thyroid lobe which obscured the surgical field. The procedure was converted to general anesthesia and exploration was expanded to the upper pole of the thyroid where an enlarged left superior parathyroid gland was identified and removed. On pathologic review the gland was hypercellular and weighed 286 mg. A post excision intact PTH level measured 15 minutes after excision remained high (89 pg/ml), and a four-gland exploration of the neck was performed. Enlarged right superior (866 mg) and inferior parathyroid glands (256 mg) were identified and removed. The left inferior parathyroid was partially resected and intraoperative pathologic evaluation identified this gland as also hypercellular. The remaining portion of the left inferior parathyroid was autotransplanted into the left sternocleidomastoid muscle. The post-operative course was uncomplicated and the patient was discharged home on day 2 with a serum calcium level of 9.1 mg/dL and a PTH level of 15.8 pg/mL.

### 3. Discussion

A causal association between chronic lithium therapy and hyperparathyroidism has been conclusively demonstrated. As noted earlier, patients on long-term lithium therapy have a 4 - 6 fold increased incidence of hyperparathyroidism compared to the general population [5,7]. Bendz et al. identified a 2.7% increased point prevalence of hyperparathyroidism among 124 patients on long-term lithium therapy (>15 years), while Mallette et al. reviewed studies involving 309 patients on chronic lithium therapy and identified 37 patients (12%) with hypercalcemia and 18 patients (16%) with elevated PTH levels [8-10]. McHenry et al. have reported that 80% of patients treated with lithium for 6 to 24 months experience a 10% increase in serum calcium levels. Christiansen et al. reported that both serum calcium and PTH levels increased by 30% in the same population, and several other authors have also reported a 10% - 60% increased prevalence of hypercalcemia and hyperparathyroidism in patients taking lithium [11-14].

LAH is associated with characteristic biochemical changes consisting of an elevated serum levels of calcium, magnesium and parathyroid hormone and decreased urinary calcium and cyclic adenosine monophosphate (CAMP) levels. These changes are similar to familial hypocalciuric hypercalcemia (FHH), but distinct from PHPT in which urinary calcium excretion is increased.
A large number of mechanisms have been proposed to account for the biochemical changes seen in LAH. (Table 3) To briefly summarize, lithium competitively antagonizes calcium sensing receptors (CaSR) and raises the threshold of serum calcium necessary to inhibit PTH secretion. This action not only increases parathyroid hormone secretion, but also exerts a multitude of systemic effects on the parathyroid glands, renal tubules and bone metabolism resulting in parathyroid gland hyperplasia and biochemical changes. Lithium is also known to accentuate adenoma formation by increasing PTH gene transcription, which may unmask hyperparathyroidism in patients with sub-clinical pre-existing adenomas once lithium is initiated.

As a consequence of the aforementioned systemic effects, it seems reasonable to conclude that lithium use should be associated with multiglandular parathyroid hyperplasia. However in most series, a solitary parathyroid adenoma was the most common LAH associated pathology. Awad et al. reported parathyroid adenomas in 14 out of 15 patients (93%) with LAH, and Carchman et al. identified adenomas in 12 out of 16 LAH patients (75%) [5,16]. Similarly, in a series of 12 LAH patients, Hundley et al. noted parathyroid adenomas in 9 patients (75%), while Bendz et al. reported adenoma in 72.5% of LAH patients (5 out of 8) [8,18]. Of note however, many authors have also described a higher incidence of multiglandular hyperplasia. Nordenstrom et al. identified multiglandular hyperplasia (MGH) in 83% of LAH patients (5 out of 6) [19]. Likewise, Jarhult et al. studied 71 patients with LAH and reported MGH in 37 patients (52%) [14]. Table 1 summarizes all reported LAH cases and demonstrates that parathyroid adenoma was the most common pathology identified in 104 out of 170 patients (61%) following parathyroidectomy [3,5,8,23-26]. (Table 1) In comparison, PHPT is associated with a single adenoma 85%, multigland hyperplasia (MGH) 15% and parathyroid carcinoma in 1% of patients [7]. It is important to point out that there is also an increased incidence rate of multigland hyperplasia (MGH) in LAH (39%, 66 of 170 patients) compared to the general population with PHPT (15%). This increased MGH incidence is likely attributable to the systemic action of lithium, and as a result may effect interpretation of localization studies (increased false negative rate, poor sensitivity) and surgical planning (increased need of bilateral neck exploration) [3,4,6,18].

The initial management of LAH is medical intervention which includes discontinuation of lithium or alternative treatment such as atypical antipsychotics and calcimimetics. (Table 2) Data relating to the efficacy of LAH medical management is available for only 10 of 194 reported LAH patients (5%). As such it is difficult to make any meaningful conclusions [12,23-26,34]. In many LAH patients, discontinuation of lithium carbonate is not medically feasible due to increased propensity of relapse [35,36]. Although serum Ca\(^{2+}\) and PTH levels tend to normalize within 4 - 6 weeks after lithium discontinuation, this change is mostly seen in patients receiving lithium for a relatively short-term defined as 5 - 10 years, or in those with only mild hypercalcemia [4]. Khandawala et al. reported on one patient who was taking lithium for 5 years, discontinuation of Li\(^+\) resulted in normocalcemia and a normal PTH level which was maintained at 5 months follow-up [24]. Dugal et al. reported on a single patient who had been on lithium therapy for 10 years, in whom discontinuation of lithium and simultaneous saline diuresis with furosemide also achi-

<table>
<thead>
<tr>
<th>Mechanism of action</th>
<th>Explanation</th>
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<tbody>
<tr>
<td>Antagonism of cations [28,29]</td>
<td>Li(^+) competitively antagonizes cations at calcium receptor at several locations—kidney, bone and parathyroid gland—causing acute hypercalcemia and hyper-magnesemia</td>
</tr>
<tr>
<td>↑ Set point or reset “calci-stat” [4,12,30, 31]</td>
<td>Li(^+) antagonizes calcium sensing receptor (CASR) causing a shift in inhibitory set point for PTH secretion, which then requires a higher levels of calcium level to block PTH secretion. This results in an overproduction of PTH despite elevated serum calcium levels</td>
</tr>
<tr>
<td>Mitogenic action [16,32,33]</td>
<td>Li(^+) exacerbates a pre-existing adenoma or PHPT—this occurs especially in patients presenting with psychiatric symptoms—the lithium therapy worsens hypercalcemia by promoting PTH overproduction. Nearly 55% of LAH have parathyroid adenomas (single or double).</td>
</tr>
<tr>
<td>↑ Parathyroid volume and parathyroid hyperplasia [2,5,11]</td>
<td>LH is associated with multiglandular hyperplasia in 25% - 55% cases (Table 1) compared to 15% in PHPT, which is attributed to direct stimulation of parathyroid gland by lithium. In patients on chronic lithium therapy neck ultrasound demonstrates increased parathyroid gland volume.</td>
</tr>
</tbody>
</table>

Abbreviations: PHPT—primary hyperparathyroidism; PTH, parathyroid hormone; Li\(^+\), lithium; CASR, calcium sensing receptor; IMP, inositol monophosphate; GSK-3b, glycogen synthase kinase-3b. Note: Several different mechanisms have been proposed by which lithium may cause hypercalcemia, ↑ PTH levels, ↑ magnesium and ↓ urinary calcium levels.
eved normocalcemia [26]. In contrast, drug withdrawal from patients on chronic lithium therapy (>10 years), has a lower rate of hypercalcemia resolution [4]. Carchman et al. noted that even after discontinuation of lithium for more than 5 months, 8 out of 16 patients on chronic lithium therapy experienced persistent hypercalcemia [16]. In LAH patients who have mild hypercalcemia or were asymptomatic, alternative medical therapy such as valproate, carbamazepine or atypical antipsychotics (risperidone) may be treatment alternatives, however at present there is very limited data to assess their efficacy in LAH patients [2,4].

Recently, a new class of drugs (calcimimetics, example: cinacalcet) has been successfully used to treat LAH with or without discontinuation of lithium [34]. (Table 3) Calcimimetics activate calcium-sensing receptors (blocked by lithium) which in turn reduce PTH secretion and prevent parathyroid hyperplasia. To date, a biochemical remission was achieved in 6 cases using cinacalcet among patients on chronic lithium therapy [12, 23,25,34]. Gregoor et al. reported resolution of LAH in 3 patients using cinacalcet while lithium was continued [34]. Szalat et al. used cinacalcet in an LAH patient who had a post-operative recurrence of hyperparathyroidism and reported resolution of hypercalcemia despite persistently elevated PTH levels [12]. Although above results are intriguing, the consistently reproducible results of parathyroidectomy have made surgical therapy the mainstay of LAH management, particularly among symptomatic patients with moderate to severe hypercalcemia, and those in whom discontinuation of Li+ is not feasible or has failed [4].

The most common indication for parathyroidectomy in LAH patients include worsening psychosomatic (50% - 60%) (fatigue, weakness, decreased concentration, short-term memory loss, depression), bone (30% - 40%) (bone pain, reduce bone mineral density, osteopenia or osteoporosis), gastrointestinal symptoms (20% - 30%) (constipation, nausea or pancreatitis), cardiac dysrhythmias and renal symptoms (1% - 5%) (decreased GFR, decreased creatinine clearance or nephrocalcinosis) [3,5,12,15,18].

Today, preoperative localization studies (ultrasound of neck, sestamibi scan and computerized tomography (CT) scan of neck) have been routinely incorporated into the management algorithm of PHPT patients, and resulted in the near universal use of focused neck exploration (90% - 95%) for this group. However, there is a paucity of data supporting the utility of preoperative localization studies in LAH patients. Among 195 LAH patients who underwent parathyroidectomy, preoperative localization studies were utilized in only 22 patients (13%). In the largest series involving 18 patients, Carchman et al. used sestamibi scan or ultrasound in 16 patients (94%). Imaging studies predicted a single adenoma in 10 patients, 2 (20%) of which had MGH at exploration [19]. In the same study, pre-operative localization studies identified MGH in 4 patients, however 2 of these patients subsequently were found to have only a single adenoma at exploration. Limited or focused neck exploration was possible in only 8 of 16 patients (50%) aided by both pre-operative localization studies and IOPTH monitoring. Szalat et al. have also reported using preoperative neck ultrasound and parathyroid scan which identified solitary uptake in 3 of 4 cases. However, at exploration IOPTH testing failed to return to baseline or decrease by >50% (positive response), and all patients required bilateral neck exploration (BNE) [12]. Despite the small number of patients included in these studies, they nevertheless highlight the poor sensitivity and limited utility of pre-operative localization studies for precise localization of pathologically enlarged parathyroid glands in LAH and in planning surgical approach [6].

Parathyroidectomy is the mainstay of treatment for LAH patients. Given the reported higher incidence of multiglandular hyperplasia (39%) and double adenomas (7%), it is not surprising that bilateral neck exploration (BNE) is the most commonly reported surgical approach in LAH. Among 170 patients with LAH, BNE was used in 162 patients (95%), whereas focused neck exploration was performed in only 8 patients (5%). In the largest reported LAH series, Jarhult et al. utilized BNE in all but 2 of 71 patients who underwent parathyroidectomy [14].

As noted earlier, intraoperative parathyroid hormone monitoring (IOPTH) has been extensively validated in patients with PHPT, and prompted more limited surgical exploration [7]. To date, only 3 studies involving LAH patients (N = 32) have assessed the usefulness of IOPTH monitoring [12,18,19]. Hundley et al. utilized IOPTH monitoring in 12 patients with LAH, of which 6 patients underwent MGH resection prompted by IOPTH monitoring [18]. In a study of 16 patients, Carchman et al. performed focused parathyroidectomy in 8 patients, while 8 patients underwent BNE guided by IOPTH monitoring [19]. Of note, these authors were able to perform a limited surgical exploration in 50% of LAH patients, compared to a 5% rate in other LAH series [12,14].

Parathyroidectomy results in biochemical remission leading to normocalcemia and normal PTH levels in the immediate post-operative period and improves psychosomatic symptoms in 90% - 97% of LAH patients. Awad et al. observed eucalcemia in 15 out of 16 patients who had parathyroidectomy for LAH; however, one patient subsequently developed recurrence at 2 years and underwent neck re-exploration with removal of an additional parathyroid adenoma [5]. At a median follow-up of 6 months (5 - 50 months range), Carchman et al. noted that
all patients were eucalcemic, however 2 patients had mild PTH elevation [19]. In contrast, Jarhult et al. reported that among 71 LAH patients who underwent parathyroidectomy, 30 patients (42%) had persistent or recurrent hyperparathyroidism at a median follow-up of 6.3 years [14]. In this series, six out of 30 patients with persistent/recurrent HPT underwent surgical re-exploration but only 2 patients were cured of hypercalcemia, one had permanent hypocalcaemia.

4. An Evidence Based Algorithm for the Management of LAH

Knowledge of the higher prevalence of hyperparathyroidism in chronic lithium users coupled with published data on the medical and surgical treatment of LAH provides a basis for developing an algorithmic approach for the treatment of LAH. (Figure 1) Serum calcium levels should be drawn before initiating lithium therapy (to rule-

Figure 1. An evidence based algorithm for the management of lithium associated hyperparathyroidism. Abbreviations: S. Ca\(^{2+}\), Serum calcium; S. Mg\(^{2+}\), Serum magnesium; S. PTH-serum parathormone; CVS-cardiovascular system; GI-gastrointestinal system; 4D-CT, 4-dimensional computed tomography; SPECT), Single Photon Emission Computed Tomography scan; IOPTH, Intra-operative parathormone monitoring; BNE, bilateral neck exploration.
out pre-existent primary HPT) and monitored within 4-6 weeks of beginning lithium therapy, and every 3 months thereafter. Serum PTH levels should be measured whenever hypercalcemia is identified. A confirmatory diagnosis of LAH necessitates ruling-out other causes of HPT. Asymptomatic patients with mild to moderate hypercalcemia may be monitored at regular intervals. The National Institute of Health (NIH) guidelines for the treatment of PHPT are also useful in LAH patients in regards to patient selection for parathyroidectomy [37]. (Figure 1) When medically feasible lithium should be discontinued or alternative therapy substituted (valproate, carbamazepine or atypical antipsychotics) in selected LAH patients. Cinaclacet hydrochloride, a calcimimetic agent, may have modest efficacy in treating LAH, and be beneficial to patients in whom discontinuation of lithium is not medically feasible or when surgery is contraindicated. In the vast majority of symptomatic patients, and those in whom either discontinuation of lithium is not possible or when alternative therapy has failed to correct hypercalcemia, parathyroidectomy is required. While preoperative localization studies may provide useful information regarding the number and location of enlarged parathyroid glands, the increased incidence of multiglandular disease in LAH, as well as a higher risk of recurrent or persistent HPT, mandates meticulous surgical exploration to visualize and remove all pathologically enlarged gland(s). Whereas a single parathyroid adenoma is found in the majority (61%) of LAH patients, limited or focused surgical exploration may be attempted when preoperative localization studies suggest feasibility, but must be further confirmed with IOPTH monitoring. A very high incidence of persistent and recurrent disease in LAH patients necessitates long-term follow-up and strict surveillance (Figure 1).

In summary, lithium use in psychiatric patients may be associated with hypercalcemia and elevated PTH levels referred to as lithium associated hyperparathyroidism (LAH). An algorithmic approach and strict post-operative surveillance protocols may improve treatment success and identify recurrent or persistent disease (Figure 1).

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