RESEARCH

Effect of autotransplantation of a parathyroid gland on hypoparathyroidism after total thyroidectomy

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Abstract

Background: The effect of parathyroid autotransplantation on hypoparathyroidism is not fully understood. The purpose of the study was to determine the effect of autotransplantation of a parathyroid gland on the incidence of hypoparathyroidism and recovery of parathyroid function at 6 months after total thyroidectomy with central neck dissection for papillary thyroid carcinoma.

Methods: All patients with autotransplantation of a parathyroid gland (no inadvertent parathyroidectomy) (group A), in situ preservation of all parathyroid glands (no autotransplantation and inadvertent parathyroidectomy) (group B) or inadvertent removal of a parathyroid gland (no autotransplantation) (group C) who underwent first-time total thyroidectomy with central neck dissection for papillary thyroid carcinoma between January 2013 and June 2016 were included retrospectively.

Results: Of the 702 patients, 383, 297 and 22 were respectively included in the groups A, B and C. The overall rates of transient and permanent hypoparathyroidism were 37.6% and 1.0%. The incidence of transient hypoparathyroidism was 43.9, 29.0 and 45.5% (A vs B, \( P = 0.000 \); A vs C, \( P = 1.000 \)), and the incidence of permanent hypoparathyroidism was 1.0, 0.7 and 4.5% (\( P > 0.05 \)). The recovery rates of serum parathyroid hormone levels were 71.4, 72.2 and 66.0% at 6-month follow-up (\( P > 0.05 \)).

Conclusion: Autotransplantation of a parathyroid gland does not affect the incidence of permanent hypoparathyroidism, but increases the risk of transient hypoparathyroidism when the rest of parathyroid glands are preserved in situ. At least 2 parathyroid glands should be preserved during total thyroidectomy with central neck dissection to prevent permanent hypoparathyroidism.

Introduction

Total thyroidectomy with central neck dissection has been adopted in many specialized endocrine surgery units for the treatment of papillary thyroid carcinoma (PTC) (1, 2, 3). However, central neck dissection, especially bilateral central neck dissection, always increases the risk of postoperative hypoparathyroidism (3). The reported incidence in the literature varies from 14 to 51.9% for transient hypoparathyroidism and 0–43% for permanent hypoparathyroidism (3, 4, 5, 6). Although transient hypoparathyroidism will recover within few months, it occasionally prolongs hospitalization or leads to readmission (4). Permanent hypoparathyroidism is a severe and potentially lethal complication. It not only increases the overall costs of thyroid surgery, but also accounts for the main category of thyroidectomy-related claims (7, 8).
Hypoparathyroidism is caused by injury to the parathyroid glands from ligation of the blood supply, inadvertent or deliberate removal, improper use of electrocauterization or destruction due to hematoma formation (9, 10). Meticulous dissection is a preferred method to preserve parathyroid glands with their blood supply in situ and to prevent inadvertent parathyroidectomy. However, parathyroid glands are still devascularized or occasionally found in the surgical specimens because of the anatomic location. Autotransplantation of devascularized or unintentionally removed parathyroid glands is generally recommended to prevent postoperative hypoparathyroidism (10, 11). Some endocrine surgeons even adopted routine autotransplantation of at least 1 parathyroid gland or all identifiable parathyroid glands to minimize the incidence of permanent hypoparathyroidism (12, 13, 14, 15). However, several studies demonstrated that parathyroid autotransplantation increased the risk of transient and permanent hypoparathyroidism (16, 17, 18). In addition, a recent study found similar incidence of permanent hypoparathyroidism whether a parathyroid gland was autotransplanted or inadvertently excised (19). Inadvertent parathyroidectomy is commonly regarded as one of the risk factors of postoperative hypoparathyroidism (20, 21). Interestingly, some studies reported that it had no relationship to the complication (22, 23, 24), suggesting that rather than removal of one or more parathyroid glands, autotransplantation renders them dysfunctional and causes postoperative hypoparathyroidism.

Therefore, the effect of parathyroid autotransplantation on postoperative hypoparathyroidism is not fully understood. In this study, we compared the patients with autotransplantation of a parathyroid gland to those with all the parathyroid glands preserved in situ and those with inadvertent removal of a parathyroid gland. The purpose of the study was to determine whether autotransplantation of a parathyroid gland affected the incidence of postoperative hypoparathyroidism and recovery of parathyroid function at 6 months after total thyroidectomy with central neck dissection for papillary thyroid carcinoma.

Methods

Patients

We retrospectively searched for all PTC patients who underwent first-time total thyroidectomy with central neck dissection (including lateral neck dissection) in the Department of Thyroid Surgery, West China Hospital of Sichuan University between January 2013 and June 2016. Those patients with autotransplantation of a parathyroid gland (no inadvertent parathyroidectomy) (group A), in situ preservation of all the parathyroid glands (no autotransplantation and inadvertent parathyroidectomy) (group B) or inadvertent removal of a parathyroid gland (no autotransplantation) (group C) were included for analyses. The patients with preoperative parathyroid dysfunction, resurgery, completion thyroidectomy, endoscopic thyroidectomy or loss of 6-month follow-up were excluded. Informed consent of at least 6-month follow-up was obtained from each patient. The study was approved by the Medical Ethics Committee of West China Hospital, Sichuan University.

Indications and surgical procedures of total thyroidectomy with lymph node dissection

In our department, the indications and surgical procedures of total thyroidectomy with lymph node dissection for PTC treatment have been described in detail previously (25, 26). Unilateral central neck dissection is performed routinely. All patients underwent therapeutic lateral neck dissection (preoperative fine-needle aspiration cytology). All surgical procedures were performed by an experienced surgeon (Zhu). Each parathyroid gland was tried to be preserved in situ intraoperatively. Thymectomy was avoided except for tumor invasion. Surgical specimens (thyroid gland and central lymphadenectomy specimen) were examined routinely for any unintentionally removed parathyroid glands. If a parathyroid gland was nonviable or resected unintentionally, it was autotransplanted into the contralateral sternocleidomastoid muscle after the confirmation by intraoperative frozen biopsy. Precise information on the number of parathyroid glands identified and autotransplanted was recorded in operative notes. A same group of experienced pathologists analyzed all surgical specimens. The presence of a whole parathyroid gland or parathyroid tissue fragments (≤1 mm) in the perithyroidal area and/or the central lymphadenectomy specimen reported by pathologists was defined as inadvertent parathyroidectomy (2).

Perioperative management

Each patient accepted preoperative assessment including serum calcium, parathyroid hormone (PTH), thyroid function, neck ultrasound and laryngoscopy. The levels of serum calcium and PTH were routinely obtained on
the first postoperative day and 1 month and 6 months after surgery. Intravenous calcium supplementation (calcium gluconate 10%, 40 mL) was routinely applied at the end of surgery. A standard dose of calcium carbonate (600–2400 mg/day) was orally administered on the first postoperative day if the serum PTH level was below the normal range. Calcitriol at 0.25–0.5 μg/day was simultaneously taken to promote absorption and reabsorption of calcium. Patients with normal PTH levels were not treated with oral calcium. Intravenous calcium was supplemented for the treatment of symptomatic hypocalcemia regardless of oral calcium and calcitriol administration. Oral calcium and calcitriol were ceased within 1 month after hospital discharge if possible. Hypoparathyroidism was defined as serum PTH level <1.6 pmol/L after surgery (normal range, 1.6–6.9 pmol/L). If serum PTH level returned to normal within 6 months postoperatively, hypoparathyroidism was classified as transient. On the contrary, it was classified as permanent if it persisted more than 6 months (27).

Data collection
The demographic characteristics, preoperative assessment, details of surgical extent (unilateral or bilateral central neck dissection), number of parathyroid glands identified, autotransplanted and inadvertently removed, final pathology, incidence of postoperative hypoparathyroidism and serum PTH levels at 6-month follow-up of all included patients were collected. Operative notes and pathology reports were reviewed independently by two authors (Su A and Gong Y) to determine the detail of parathyroid glands. PTC was staged by the system of American Joint Committee on Cancer (AJCC) (the 7th edition) (28). The primary endpoints were the number of parathyroid glands identified, autotransplanted and inadvertently removed, incidence of postoperative hypoparathyroidism and serum PTH levels at 6-month follow-up.

Statistical analysis
Continuous variables were expressed as mean ± S.D. The data were analyzed by SPSS software (version 19.0, SPSS, Inc., 1989–2010). Statistical comparison between groups A and B or groups A and C was calculated by the Fisher’s exact test or Student’s t-test. Multivariate analyses were performed to assess the independent risk factors of transient and permanent hyperparathyroidism. For multivariate analyses, the variables that were statistically significant in univariate analyses were included in logistic regression analyses. The results of the multivariate analyses were expressed as odds ratio (OR) and 95% confidence interval (CI). Statistical significance was set at P < 0.05.

Results
Patient characteristics
Of the 1492 patients whose medical records were reviewed, 702 met the study criteria and were included in the study (Fig. 1). There were 192 male and 510 female, with a mean age of 42.6 ± 12.9 years. Unilateral central neck dissection and bilateral central neck dissection were performed on 253 and 356 patients, respectively. Seventy-seven patients underwent bilateral central neck dissection with unilateral neck dissection, and 16 accepted bilateral central neck dissection with bilateral neck dissection. Groups A, B and C consisted of 383, 297 and 22 patients, respectively. Table 1 shows a comparison of baseline characteristics of the patients in the 3 groups. No significant differences were found between groups A and B or groups A and C in age, gender, body mass index (BMI), comorbidities (hypertension, diabetes, thyroiditis, Graves’ disease, hypothyroidism and nodular goiter) and preoperative levels of serum calcium and PTH. Patients with autotransplantation of a parathyroid gland underwent more bilateral central neck dissection than those with preservation of all the parathyroid glands in situ (67.4% vs 58.6%, P = 0.020).

Characteristics of tumors and central lymph nodes
Table 2 shows the characteristics of tumors and central lymph nodes. There were no significant differences between groups A and B or groups A and C in the largest tumor size, multifocality, primary tumor location, T classification, N classification, AJCC stage and gross extrathyroidal extension. Significantly more central lymph nodes and metastatic central lymph nodes were retrieved in the group A than in the group B (P = 0.001 and P = 0.024, respectively). However, there was no significant difference of the two parameters between groups A and C (P = 0.731 and P = 0.368, respectively).

Details of parathyroid glands, hypoparathyroidism and 6-month follow-up
Among the 3 groups, less parathyroid glands were identified intraoperatively in the patients with inadvertent removal
of a parathyroid gland. Rates of transient and permanent hypoparathyroidism for the entire cohort were 37.6% (264/702) and 1.0% (7/702), respectively. The incidence of transient hypoparathyroidism was 43.9% (168/383) in patients with autotransplantation of a parathyroid gland, 29.0% (86/297) in those with preservation of all the parathyroid glands and 45.5% (10/22) in those with inadvertent removal of a parathyroid gland (43.9% vs 29.0%, \( P = 0.000 \) and 43.9% vs 45.5%, \( P = 1.000 \)). The incidence of permanent hypoparathyroidism in the 3 groups was 1.0, 0.7 and 4.5%, respectively (\( P > 0.05 \)).

Table 3 shows the serum PTH levels in the 3 groups at 6-month follow-up. Compared to the patients with in situ preservation of all the parathyroid glands, significantly lower serum PTH level was found in patients with autotransplantation of a parathyroid gland on
postoperative day 1. However, similar levels of serum PTH were found in the 3 groups after 1 month and 6 months ($P > 0.05$). After 6 months of follow-up, the recovery rates of serum PTH level were 71.4, 72.2 and 66.0% in the 3 groups.

**Risk factors for transient and permanent hypoparathyroidism**

Perioperative factors, which influenced the incidence of transient and permanent hypoparathyroidism, were sought by univariate and multivariate analyses. Gender, surgical extent and parathyroid autotransplantation had significant effects on the formation of transient hypoparathyroidism and the number of parathyroid glands preserved in situ was associated with permanent hypoparathyroidism on univariate analyses (Table 4). Multivariate analysis found female, bilateral central neck dissection and autotransplantation of a parathyroid gland to be independent risk factors of transient hypoparathyroidism (Table 5). Preservation of less than 2 parathyroid glands was an independent risk factor of permanent hypoparathyroidism (OR, 9.085; 95% CI, 1.981–44.660; $P = 0.005$).

**Discussion**

Parathyroid autotransplantation, first described in humans by Lahey in 1926 (29), is regarded as a major technical and physiological breakthrough in the field of thyroid surgery. In recent years, it has been increasingly employed to prevent postoperative hypoparathyroidism.

**Table 3** The details of parathyroid glands, hypoparathyroidism and the results of follow-up in the 3 groups.

<table>
<thead>
<tr>
<th>Variables</th>
<th>Group A ($n=383$)</th>
<th>Group B ($n=297$)</th>
<th>Group C ($n=22$)</th>
<th>$P$ value*</th>
<th>$P$ value†</th>
</tr>
</thead>
<tbody>
<tr>
<td>Parathyroid identification</td>
<td>3.5 ± 0.8</td>
<td>3.4 ± 0.8</td>
<td>2.7 ± 0.8</td>
<td>0.677</td>
<td>0.000</td>
</tr>
<tr>
<td>Hypoparathyroidism</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Transient</td>
<td>168</td>
<td>86</td>
<td>10</td>
<td>0.000</td>
<td>1</td>
</tr>
<tr>
<td>Permanent</td>
<td>4</td>
<td>2</td>
<td>1</td>
<td>0.701</td>
<td>0.245</td>
</tr>
<tr>
<td>Postoperative PTH (pmol/L)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1 day</td>
<td>1.76 ± 1.18</td>
<td>2.34 ± 1.41</td>
<td>1.93 ± 1.34</td>
<td>0.001</td>
<td>0.664</td>
</tr>
<tr>
<td>1 month</td>
<td>3.45 ± 1.63</td>
<td>3.74 ± 1.70</td>
<td>3.50 ± 1.65</td>
<td>0.220</td>
<td>0.920</td>
</tr>
<tr>
<td>6 months</td>
<td>3.87 ± 1.61</td>
<td>4.08 ± 1.61</td>
<td>3.79 ± 1.63</td>
<td>0.372</td>
<td>0.879</td>
</tr>
</tbody>
</table>

*Groups A vs B; †groups A vs C.

PTH, parathyroid hormone.
Table 4  Univariate analysis of risk factors for the development of transient and permanent hypoparathyroidism.

<table>
<thead>
<tr>
<th>Variables</th>
<th>No permanent</th>
<th>Permanent</th>
<th>P value †</th>
<th>P value ‡</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Normal (n=431)</td>
<td>Transient (n=264)</td>
<td>Total (n=695)</td>
<td></td>
</tr>
<tr>
<td>Age (&lt;45/≥45, years)</td>
<td>241/190</td>
<td>152/112</td>
<td>393/302</td>
<td>2/5</td>
</tr>
<tr>
<td>Gender (male/female)</td>
<td>138/293</td>
<td>53/211</td>
<td>191/504</td>
<td>1/6</td>
</tr>
<tr>
<td>BMI (&lt;25/≥25, kg/m²)</td>
<td>303/128</td>
<td>195/69</td>
<td>498/197</td>
<td>5/2</td>
</tr>
<tr>
<td>Hypertension</td>
<td>57</td>
<td>26</td>
<td>83</td>
<td>1</td>
</tr>
<tr>
<td>Diabetes</td>
<td>12</td>
<td>6</td>
<td>18</td>
<td>0</td>
</tr>
<tr>
<td>Thyroiditis</td>
<td>121</td>
<td>78</td>
<td>199</td>
<td>2</td>
</tr>
<tr>
<td>Graves’ disease</td>
<td>12</td>
<td>8</td>
<td>20</td>
<td>0</td>
</tr>
<tr>
<td>Hypothyroidism</td>
<td>5</td>
<td>3</td>
<td>8</td>
<td>0</td>
</tr>
<tr>
<td>Nodular goiter</td>
<td>251</td>
<td>158</td>
<td>409</td>
<td>3</td>
</tr>
<tr>
<td>Preoperative calcium (mmol/L)</td>
<td>2.31±0.16</td>
<td>2.32±0.44</td>
<td>2.31±0.30</td>
<td>2.26±0.09</td>
</tr>
<tr>
<td>Preoperative PTH (pmol/L)</td>
<td>5.73±1.98</td>
<td>5.28±1.75</td>
<td>5.54±1.16</td>
<td>4.28±2.33</td>
</tr>
<tr>
<td>Surgical extent (UCND/BCND)</td>
<td>172/259</td>
<td>79/185</td>
<td>251/444</td>
<td>2/5</td>
</tr>
<tr>
<td>Largest tumor size (mm)</td>
<td>13.6±8.4</td>
<td>14.6±10.3</td>
<td>14.0±9.2</td>
<td>14.4±5.4</td>
</tr>
<tr>
<td>Multifocality</td>
<td>101</td>
<td>76</td>
<td>177</td>
<td>2</td>
</tr>
<tr>
<td>Tumor location (upper/middle/lower/isthmus)</td>
<td>127/174/161/164</td>
<td>64/121/70/9</td>
<td>191/295/186/23</td>
<td>4/3/0/0</td>
</tr>
<tr>
<td>T classification (T1–2/T3–4)</td>
<td>175/256</td>
<td>118/146</td>
<td>293/402</td>
<td>3/4</td>
</tr>
<tr>
<td>N classification (NO/N1)</td>
<td>166/263</td>
<td>94/170</td>
<td>260/435</td>
<td>2/5</td>
</tr>
<tr>
<td>AJCC stage (I–II/III–IV)</td>
<td>272/159</td>
<td>179/85</td>
<td>451/244</td>
<td>4/3</td>
</tr>
<tr>
<td>Gross extrathyroidal extension</td>
<td>174</td>
<td>106</td>
<td>280</td>
<td>3</td>
</tr>
<tr>
<td>Harvested lymph nodes (&lt;10/≥10)*</td>
<td>205/226</td>
<td>107/157</td>
<td>312/383</td>
<td>5/2</td>
</tr>
<tr>
<td>Parathyroid identification (&lt;3/≥3)</td>
<td>55/376</td>
<td>39/225</td>
<td>94/601</td>
<td>3/4</td>
</tr>
<tr>
<td>Parathyroid autotransplantation (0/1)</td>
<td>220/211</td>
<td>96/168</td>
<td>316/379</td>
<td>3/4</td>
</tr>
<tr>
<td>Inadvertent parathyroidectomy (0/1)</td>
<td>420/01</td>
<td>254/10</td>
<td>674/21</td>
<td>6/1</td>
</tr>
<tr>
<td>Parathyroid preservation (&lt;2/≥2)</td>
<td>29/402</td>
<td>23/241</td>
<td>52/643</td>
<td>3/4</td>
</tr>
</tbody>
</table>

*Central lymph nodes; †normal vs transient; ‡no permanent (total) vs permanent.

AJCC, American Joint Committee on Cancer; BCND, bilateral central neck dissection; BMI, body mass index; PTH, parathyroid hormone; UCND, unilateral central neck dissection.
Table 5  Multivariate analysis of risk factors for the development of transient hypoparathyroidism.

<table>
<thead>
<tr>
<th>Variables</th>
<th>OR</th>
<th>95% CI</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gender (female)</td>
<td>1.977</td>
<td>1.366–2.861</td>
<td>0.000</td>
</tr>
<tr>
<td>Surgical extent (≥BCND)</td>
<td>1.554</td>
<td>1.114–2.168</td>
<td>0.009</td>
</tr>
<tr>
<td>Parathyroid autotransplantation (1)</td>
<td>1.834</td>
<td>1.332–2.526</td>
<td>0.000</td>
</tr>
</tbody>
</table>

BCND, bilateral central neck dissection; CI, confidence interval; OR, odds ratio. (1) autotransplantation of one parathyroid gland increases the incidence of transient hypoparathyroidism.

(10, 11, 12, 13, 14, 15). However, there is no good evidence to prove that parathyroid grafts do in fact prevent this complication because the recovery of parathyroid function depends on both preserved and autotransplanted parathyroid glands (19). As shown in the present study, there is large consensus that autotransplantation increases the risks of transient hypoparathyroidism and postoperative hypocalcemia (8, 30, 31). It is still controversial whether it can reduce the incidence of permanent hypoparathyroidism. Although two studies reported that autotransplantation of all identifiable parathyroid glands could lower the risk of permanent hypoparathyroidism (14, 15), Kihara and coworkers (17) found that the strategy led to a higher permanent hypoparathyroidism rate. The effect of autotransplantation on permanent hypoparathyroidism is closely related to the number of autotransplanted parathyroid glands (8, 17). Kikumori and coworkers (15) revealed that the incidence of permanent hypoparathyroidism was inversely correlated with the number of autotransplanted parathyroid glands. Several studies reported that autotransplantation of at least one parathyroid gland effectively decreased the incidence of permanent hypoparathyroidism (12, 13, 30). A multicenter study also investigated the impact of autotransplantation of a parathyroid gland on postoperative hypoparathyroidism and drew a similar conclusion (32). These conclusions, however, are questioned by the confounding factors, including different number of parathyroid glands autotransplanted and preserved, a variety of thyroid gland disorders (benign and malignant tumors) and different surgical times (first-time surgery and resurgery).

In addition, these studies did not provide the comparison with inadvertent parathyroidectomy and the change of serum PTH levels after follow-up.

There is no doubt that parathyroid autotransplantation is an effective procedure to preserve parathyroid function. Direct evidence of grafts function can be obtained through the comparison of serum PTH levels between the transplanted and non-transplanted arms (33, 34). The reported success rate of autotransplantation ranges from 55 to 100% (5). In our study, although there was no significant difference of the serum PTH levels among the 3 groups at 6-month follow-up, the recovery rates were higher in patients with autotransplantation of a parathyroid gland or in situ preservation of all the parathyroid glands than those with inadvertent removal of a parathyroid gland (71.4%/72.2% vs 66.0%). The parathyroid function of the patients without inadvertent parathyroidectomy in the 2 groups did not recover to 100% of the preoperative levels at 6-month follow-up, which is attributed to the thermal and/or mechanical injury of parathyroid glands, dysfunction of parathyroid glands preserved in situ, loss of part of parathyroid tissue for intraoperative frozen biopsy, dysfunction of the grafts and/or short-term follow-up period.

Most available reports on parathyroid autotransplantation have overestimated its impact and underestimated the effect of in situ preservation on parathyroid function (12, 13, 30, 32). In our study, the prevalence of permanent hypoparathyroidism after total thyroidectomy with central lymph node dissection was similar whether a parathyroid gland was autotransplanted, preserved or inadvertently removed when the rest of parathyroid glands were preserved in situ. This observation implies that the part of function preserved by autotransplantation of a parathyroid gland has no significant effect on the incidence of permanent hypoparathyroidism. Therefore, parathyroid glands preserved in situ, instead of the one autotransplanted, are critical in preventing permanent hypoparathyroidism (2). This finding can be explained by the fact that preserving at least 2 parathyroid glands with an intact blood supply is sufficient to prevent permanent hypoparathyroidism (35).

The current study also demonstrated that in situ preservation of less than 2 parathyroid glands was a significant variable favoring permanent hypoparathyroidism. Therefore, at least 2 parathyroid glands should be carefully preserved during total thyroidectomy with central neck dissection to avoid permanent hypoparathyroidism.

Intraoperative identification of parathyroid glands is negatively correlated with inadvertent removal of parathyroid glands (36). Inadvertent parathyroidectomy is always reported to add a further risk of postoperative transient and permanent hypoparathyroidism (20, 21, 37). A recent study revealed that presence of at least 2 parathyroid glands in surgical specimens was a risk factor of transient hypoparathyroidism (38). However, one study revealed no correlation between inadvertent
Autotransplantation of a parathyroid gland

Declaration of interest
The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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Author contribution statement
Anping Su helped in writing and revising the manuscript. Yanping Gong, Wenshuang Wu, Rixiang Gong and Zhihui Li helped in revising the manuscript. Jinqiang Zhu helped in revising the manuscript and is a corresponding author.

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