Continuous, Large-Volume Hydrodissection to Protect Delicate Structures around the Thyroid throughout the Radiofrequency Ablation Procedure

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Keywords

Hydrodissection · Benign thyroid nodules · Radiofrequency ablation · Ultrasound-guided ablation · Interventional ultrasonography

Abstract

Introduction: Small-volume hydrodissection liquid dissipates rapidly and confers only short-term protection during radiofrequency ablation (RFA) of benign thyroid nodules. The aim of this study was to establish a safe method for continuous, large-volume hydrodissection. **Methods:** A long needle was inserted and positioned outside the thyroid capsule; 5% glucose was injected to maintain a 3- to 5-mm continuous safety buffer. From October 2015 to July 2020, 166 patients underwent hydrodissection with different volumes, and ablation efficacy and complications associated with different liquid volumes (≤ 40 mL vs. > 40 mL) were compared at 1-month postprocedure. Moreover, 20 mL liquid (equivalent to 250 mL in the human body) was injected around the thyroid of a rhesus monkey, after which CT scans were used to visualize the liquid's fate and verify its safety. **Results:** The 51

patients with 10–40 mL injections and 116 patients with larger injections (45–450 mL) showed similar complete ablation rates (88.46% vs. 90.44%, $p = 0.582$), comparable 6-month VRR (82.79% vs. 76.62%, $p = 0.079$), and complication incidences, although the latter group had larger nodules (9.11 mL vs. 13.79 mL, $p = 0.003$), more energy delivered (3.44 kcal vs. 6.04 kcal, $p < 0.001$), and longer operation times (51.37 min vs. 69.2 min, $p < 0.001$). In the animal experiment, the 20 mL of liquid diffused quickly (within 10 min) from the vicinity of the thyroid to the mediastinum and retropharyngeal space. It was observed in the kidneys at 10 min and disappeared from the neck and chest space by 24 h. **Conclusions:** Continuous, large-volume hydrodissection can protect the delicate structures around the thyroid throughout the RFA procedure and might be beneficial in large thyroid nodule ablation.

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Introduction

Thyroid nodules constitute a frequently encountered clinical problem [1, 2], and some benign thyroid nodules (BTNs) may require treatment for associated symptoms and/or because of cosmetic problems. As curative surgery has several drawbacks [3, 4], radiofrequency ablation (RFA), a minimally invasive treatment modality, was introduced and has been reported to be safe and effective for treating BTNs [5–8]. However, the delicate structures around the thyroid are likely to be damaged during thermal ablation [9]. How to protect vital structures from heat injuries plays a significant role in RFA for BTNs.

The hydrodissection technique is a protective measure used to reduce heat injury to adjacent structures [10] and has been cited in some guidelines [11–13]. Hydrodissection involves injecting liquids around the thyroid capsule, ensuring a safe distance between the thyroid and critical structures [10–13].

However, little attention has been given to the amount of hydrodissection liquid. In a few studies, 10–40 mL was considered effective [12, 14, 15]. Nonetheless, the duration of 10–20 mL saline was only 5–15 min [16], which limits its use to short procedures. For time-consuming procedures, which can even take up to 1 h in microwave ablation [17], this duration may be too short to cover the whole ablation process. One possible solution would be to increase the volume of injected liquid. Lee M et al. [18] injected as much as 260 mL cold 5% glucose into the tracheoesophageal groove of a 47-year-old male to reduce nerve damage during RFA. Similarly, Cui et al. [19] used 200 mL saline during microwave ablation of a 5.4-cm substernal goiter. This innovative and seminal work pioneered a new approach: large-volume hydrodissection. However, no study has adequately explored whether some side effects including airway compression increase with injections exceeding 40 mL of liquid, and the location and distribution of the dwindling liquid remain poorly understood.

Based on the clinical experiences of >200 patients who underwent RFA of BTNs, we established a method of intermittent and repeated injections of hydrodissection liquid that could inject >40 mL of liquid. The complications and ablation efficacy in 166 patients who underwent this method were retrospectively analyzed. In addition, the fate of the hydrodissection liquid was verified by injecting 5% glucose around the thyroid of an adult rhesus monkey. This investigation enhances our understanding of the hydrodissection technique and might help improve the safety of time-consuming ablations.

Fig. 1. Flowchart of patient enrollment. RFA, radiofrequency ablation.

Materials and Methods

Patients

This study was approved by the Review Board of the Third Affiliated Hospital of Sun Yat-sen University and was registered at www.chictr.org.cn (ChiCTR-INR-16007884). From October 2015 to July 2020, 217 consecutive patients were treated with RFA of BTNs in our hospital. All patients signed an informed consent form before receiving treatment.

All enrolled patients fulfilled the criteria established by the Korean Society of Thyroid Radiology for BTNs [11, 12, 20] and underwent hydrodissection. The exclusion criteria were the inability to tolerate treatment and lack of postoperative 1-month follow-up information (online suppl. Table 1; see online Supplementary Materials).

After excluding 51 patients according to the above criteria, 166 patients were enrolled (Fig. 1). To explore whether there were differences in ablation efficacy and complications in patients who received different hydrodissection volumes, we divided the patients into 2 groups according to the conventional volume (40 mL) in the previous study: group 1 (hydrodissection volume ≤ 40 mL) and group 2 (hydrodissection volume > 40 mL).

Pretreatment Evaluation

Before the procedure, the ultrasonography (US) findings, fine-needle aspiration findings, contrast-enhanced US (CEUS) findings, and laboratory and clinical results were evaluated. The US examination included characterization of the location, shape, size, margins, solid/cystic proportions, echogenicity, calcification status, and internal vascularity of each nodule. The nodule volume was calculated as follows: volume = $\pi abc/6$ (a, b, and c: 3 diameters of the nodule). Laboratory tests included serum thyroid function tests and antibody tests.

Hydrodissection Technique and RFA Procedure

Patients were placed in the supine position with their necks extended. Local anesthesia with 2% lidocaine was applied from the skin puncture point to the outer capsule of the thyroid. A long

needle (22 gauge, 70-mm length; Hakko Co., Ltd.) was inserted and maintained outside the thyroid capsule, and 5% glucose was injected by using a catheter until successful hydrodissection was established. Successful hydrodissection was a 3- to 5-mm barrier between the outer thyroid capsule and common carotid artery, the posterior thyroid capsule and trachea/esophagus, the thyroid and parathyroid, and the thyroid posterior capsule and recurrent laryngeal nerves [7, 8, 12]. If no breathing difficulty was reported during injection, we began ablation. During ablation, we ensured that the tip of the long needle was located between the posterior thyroid capsule and longus colli muscle (online suppl. Figure). If the distance was <3 mm, a repeat injection was performed to guarantee a safety buffer of at least 3 mm.

We used an RF generator (VIVA RF System®; STARmed) and an internally cooled 18-gauge, 70-mm length, 7- or 10-mm, active-tip electrode (Star RF Electrode®; STARmed). The moving-shot technique was used to perform multiple ablations until the nodule was completely covered by the transient hyperechoic area [21, 22]. CEUS was used to evaluate whether the appearance of an entire nodule changed to nonenhancement. For enhanced nodules, an additional ablation was performed.

Any specific complaints or concerns were monitored immediately after the procedure and during the follow-up period. Safety was defined as the incidence of complications and side effects. Major and minor complications and side effects were defined according to the criteria established by the Society of Interventional Radiology [9, 23–26].

Follow-Up

Postprocedural follow-up was performed 1, 3, and 6 months after treatment. At the visit, a US examination, CEUS examination, and serum thyroid tests were performed. Complete ablation was defined as the entire nodule being nonenhanced on CEUS. The volume reduction ratio (VRR) and complete ablation rate (CAR) are important indicators reflecting ablation efficacy [27]:

$$\text{VRR (\%)} = [\text{initial volume (mL)} - \text{final volume (mL)}] / \text{initial volume (mL)} \times 100\% \text{ (8)},$$

$$\text{CAR} = \text{number of nodules that achieved complete ablation (N)} / \text{total number of nodule (N)} \times 100\%.$$

Animal Data

The adult male rhesus monkey was 22 years old, with a body weight of 10 kg. After general anesthesia was administered, the monkey was laid in a supine position with the neck hyperextended. Under US guidance, a 22-gauge needle was inserted outside the thyroid capsule. Then, a 20 mL mixture containing 10 mL 5% glucose and 10 mL iohexol was injected for hydrodissection. Twenty milliliters in the monkey is equivalent to 250 mL in a 60-kg human body [28]. Successful hydrodissection was defined as detailed above. A thin-layer CT scan of the neck, chest, and abdomen was performed 10 min, 1 h, 2 h, and 24 h after the liquid injection. Enhancement of the perithyroidal space, mediastinum, blood vessels, and urinary system was observed to confirm where the injected liquid went. After the monkey woke up, its vital signs were monitored for 72 h.

Statistical Analysis

Continuous parameters are expressed as mean $\pm$ standard deviation, while categorical variables are expressed as numbers and percentages. Independent *t* tests and Mann-Whitney U tests for

continuous variables and Fisher's exact tests/ χ^2 tests for categorical variables were employed to perform statistical analysis of between-group comparisons. In addition, statistical significance was set at $p < 0.05$. SPSS version 23.0 (IBM, Armonk, NY, USA) was used for all statistical analyses.

Results

A total of 166 patients with 187 nodules completed their treatment with an average hydrodissection liquid volume of 113.68 mL (10–450 mL). Among them, only 1 patient experienced hoarseness, 2 patients had hematomas, and 3 patients vomited (Fig. 2; online suppl. Table 2).

Patient age, sex, location of nodules, and serum thyroid tests were comparable between group 1 (hydrodissection volume ≤ 40 mL) and group 2 (hydrodissection volume > 40 mL) (Table 1). The maximum nodule diameter and baseline volume, solid portion, operation time, and total energy delivered in group 2 were significantly greater than those in group 1. Furthermore, group 2 had more nodules with a maximum diameter of > 40 mm than group 1 (68/135 vs. 14/52).

Regarding ablation efficacy, the CAR at the 1-month follow-up and VRR at the 6-month follow-up showed no significant difference between the 2 groups (90.44% vs. 88.46%, $p = 0.582$; $76.62\% \pm 17.74\%$ vs. $82.79\% \pm 13.78\%$, $p = 0.079$; respectively), although the VRR at the 1- and 3-month follow-ups of group 2 was lower than that of group 1 ($39.94\% \pm 28.62\%$ vs. $50.25\% \pm 23.78\%$, $p = 0.035$; $64.67\% \pm 20.50\%$ vs. $73.56\% \pm 17.63\%$, $p = 0.020$; respectively). There were no significant differences in 1-month serum tests between the 2 groups.

Regarding complications and side effects, the 2 groups did not differ significantly in major complications, minor complications, or side effects (Table 2). The number of patients who suffered pain, chest tightness, edema, and feelings of choking when eating in group 2 was clearly greater than that in group 1.

After injecting a 20 mL mixture around the thyroid of the rhesus monkey, serial CT and reconstruction images of the neck and chest showed the location and movement of the hydrodissection liquid. The liquid was injected into the visceral space first and then quickly spread around the thyroid and formed a working barrier, occupying the potential space around the neck, up to the chin, down to the mediastinum, and posterior to the retropharyngeal space in 10 min. In the next 2 h, the liquid spread more widely. The dynamic changes indicated that a potentially large space around the neck and mediastinum was able to store a large volume of liquid.

Fig. 2. A case of intermittent, repeated injection of 350 mL hydrodissection liquid. A 30-year-old female patient had a nodule in the right lobe of the thyroid. Preoperative fine-needle aspiration cytologic examination confirmed its benignancy. The blue line in the figure shows the rim of the nodule boundary. Preoperative images show that the rim of the 52 × 36 × 32 mm nodule was clear and smooth (a). Preoperative assessment of dangerous structures around the nodules. The nodule was adjacent to the cervical sympathetic nerve node, recurrent laryngeal nerve, trachea, and esophagus, and the yellow line shows the posterior cervical sympathetic ganglion (b). Glucose (5%) was injected to separate the nodule and

the surrounding structures at a distance of >3 mm (c). After assurance of a sufficient safety buffer, multiple ablations were performed until the nodule was completely covered by a transient hyperechoic area in three-dimensional space. Then, CEUS was performed again to confirm that the nodule had been completely ablated (d). One-day US images showed that the hydrodissection liquid was absorbed completely (e). One-month US images showed that the nodule achieved complete ablation and was reduced to 44 × 32 × 10 mm (VRR = 56.5%) (f). CEUS, contrast-enhanced ultrasonography; US, ultrasonography; VRR, volume reduction ratio.

Table 1. Patient characteristics and treatment parameters of 2 groups

Variables	Group 1 (hydrodissection volume ≤40 mL)	Group 2 (hydrodissection volume >40 mL)	<i>p</i> value
Patient characteristics			
Patients, <i>n</i>	51	115	
Age, years	40.67±15.53	42.89±13.29	0.348
Female/male	37/14	96/29	0.104
Nodule characteristics			
Nodules, <i>n</i>	52	135	
Location of nodules			
Left	21	59	0.641
Right	28	72	
Isthmus	3	4	
Maximum diameter, mm	32.54±8.41	38.38±10.61	<0.001
20–29	21	33	
30–39	17	34	
>40	14	68	
Baseline volume, mL	9.11±8.55	13.79±11.53	0.003
Solid, %	52.40±38.07	68.2±35.03	0.020
Baseline thyroid tests			
TSH, uIU/mL	1.63±0.99	1.22±0.80	0.008
FT3, pmol/L	4.91±3.63	4.46±0.77	0.615
FT4, pmol/L	13.83±2.63	13.50±3.13	0.373
TPOAb, IU/mL	0.41 (0.15–2.80)	0.62 (0.27–9.32)	0.098
TgAb, IU/mL	1.67 (1.04–3.05)	1.76 (1.07–11.21)	0.422
Treatment parameters			
Hydrodissection volume, mL	26.19±9.62	152.36±102.90	<0.001
Operation time, min	51.37±20.15	69.23±27.46	<0.001
Total energy delivered, kcal	3.44±2.54	6.04±4.03	<0.001
1-month follow-up			
CAR, %	88.46 (46/52)	90.44 (123/135)	0.582
Volume of nodules, mL	4.25±4.18 (10–40)	7.52±6.98 (45–450)	<0.001
VRR, %	50.25±23.78	39.94±28.62	0.035
1-month thyroid tests			
TSH, uIU/mL	1.90±3.90	1.27±0.92	0.515
FT3, pmol/L	4.52±0.91	4.53±1.49	0.698
FT4, pmol/L	14.01±2.77	13.34±2.84	0.052
TPOAb, IU/mL	0.71 (0.27–2.28)	0.61 (0.28–8.34)	0.518
TgAb, IU/mL	1.50 (1.00–2.08)	2.54 (1.13–12.53)	0.112
3-month follow-up			
Volume of nodules, mL	2.13±1.96	4.83±5.27	<0.001
VRR, %	73.56±17.63	64.67±20.50	0.020
6-month follow-up			
Volume of nodules, mL	2.00±3.44	3.09±4.05	0.032
VRR, %	82.79±13.78	76.62±17.74	0.079

Nodule volume (mL) = $\pi abc/6$ (a, b, and c: 3 diameters of the nodule). CAR: number of nodules achieved complete ablation (*N*)/total number of nodules (*N*) × 100%. VRR: (initial volume [mL] – final volume [mL])/initial volume [mL] × 100. VRR, volume reduction ratio; CAR, complete ablation rate.

Table 2. Complications and side effects of 2 groups

Complication or side effect	Group 1 (hydrodissection volume ≤40 mL)	Group 2 (hydrodissection volume >40 mL)	<i>p</i> value
Major, <i>n</i> (%)			
Voice change (>1 month)	1 (1.96)	0	0.307
Minor, <i>n</i> (%)			
Hematoma	0	2 (1.74)	1.000
Nausea and vomiting	2 (3.92)	1 (0.87)	0.465
Side effect, <i>n</i> (%)			
Pain (VAS ≥7)	3 (5.88)	8 (6.96)	1.000
Vasovagal reaction	3 (5.88)	2 (1.74)	0.343
BP change	3 (5.88)	3 (2.61)	0.554
Swelling of neck	1 (1.96)	2 (1.74)	1.000
Chest tightness	0	2 (1.74)	1.000
Edema	2 (3.92)	5 (4.35)	1.000
Coughing	1 (1.96)	2 (1.74)	1.000
Paraesthesia pharynges	5 (9.80)	8 (6.96)	0.529
Feeling of choking when eating	1 (1.96)	7 (6.09)	0.452
Dizziness	1 (1.96)	3 (2.61)	1.000
Burning sensation	3 (5.88)	1 (0.87)	0.163

Notably, the amount of liquid in the neck and chest images decreased from 10 min to 2 h, and the 24-h images showed no liquid, which indicated rapid absorption of the liquid. Moreover, there was enhancement in the kidneys, ureters, and bladder in the 10-min, 1-h, 2-h and 24-h images. The images revealed that the abundant capillaries and lymphatic vessels around the neck and mediastinum could quickly absorb the retained liquid, and the excretion route of the hydrodissection liquid was through the urinary system (Fig. 3, 4).

Regarding clinical manifestations, the monkey was continuously monitored and exhibited stable vital signs. It woke up 1 h after the last CT examination and survived for >1 month subsequently. Notably, the trachea moved to the right side slightly, without lumen diameter changes or manifestations of dyspnea. The full lumen of the trachea allowed sufficient air to be inhaled, thus avoiding serious breathing difficulties due to the injection of hydrodissection liquid.

Discussion

The hydrodissection technique is a protective measure to reduce heat injury to adjacent structures in RFA of BTNs. To maintain continuous hydrodissection protection throughout some time-consuming ablation processes, we established a method of intermittent, repeated injections of hydrodissection liquid. This method was proven to be safe and effective from bench to bed.

Since we established the method, 166 patients have been injected with an average of 113.68 mL of 5% glucose. The successful use of this large-volume hydrodissection liquid volume indicates the feasibility and safety of this proposed technique. Compared with previous studies, our method was able to apply a larger volume of liquid around the thyroid capsule [12, 14, 15]. In addition, a large volume of liquid (45~450 mL) did not increase the occurrence of complications compared with previous studies (0–10%) [25]. Regarding liquid volume, patients receiving >40 mL of hydrodissection liquid had larger BTNs, more energy delivered, and longer operation times than those receiving <40 mL. However, CAR, 6-month VRR, and incidence of complications were comparable between the 2 groups. These results indicated that a large volume of liquid was required to protect the surrounding structures because large BTNs need longer operation times and more energy delivered. Furthermore, with good protection from large-volume hydrodissection, large BTNs could be completely ablated during the first ablation, and a comparable volume reduction was achieved without increasing the incidence of complications. Our user-friendly method showed its superiority in RFA of large BTNs.

Another finding worth noting was the number of patients with side effects such as pain and chest tightness, which might be related to the compression caused by the larger volume of liquid. Indeed, more patients suffered side effects among those who received more hydrodissec-

Fig. 3. CT and reconstruction images of the 22-year-old rhesus monkey (weight = 10 kg) after injecting 20 mL hydrodissection liquid (10 mL 5% glucose and 10 mL iohexol) around the left thyroid lobe. a: CT images of the neck. a1–a5 show that 10 min to 2 h after injection, liquid spread around the visceral space and extended to the retropharyngeal space and the opposite side. Twenty-four hours after liquid injection, no liquid remained in the neck. b: schematic diagrams of the neck corresponding to a1–a5. c: recon-

struction images of the chest. c1–c5 show that 10 min to 2 h after injection, liquid spread around the visceral space and extended to the mediastinum. Twenty-four hours after liquid injection, no liquid remained in the chest. d: CT images of the abdomen. d1–d5 show that 10 min to 2 h after injection, the kidneys, ureters, and bladder were enhanced by hydrodissection liquid absorption. Twenty-four hours after liquid injection, the bladder was still enhanced as a result of residue iohexol.

tion liquid injections. In addition, although the nodule VRR of patients receiving more hydrodissection liquid was lower at the 1- and 3-month follow-up, the 6-month VRR was not significantly different between the 2 groups. This result could be explained by the fact that large BTN's need more time to absorb necrotic tissue; thus, the volume reduction in large BTN's was not as good as that in small BTN's. This is consistent with previous literature reports [7, 15, 29, 30].

Finally, we injected a 20 mL mixture around the thyroid of a rhesus monkey, which has neck structures similar to those of humans, to show the fate of the hydrodissection liquid. There were several striking findings in this experiment. First, the 20 mL mixture diffused quickly from the vicinity of the thyroid to the mediastinum and retropharyngeal space and was then cleared by the hemolymphatic system and excreted by the urinary system. This hydrodissection liquid diffusion process was similar to the spreading process of abscesses and effusions of the

Fig. 4. Schematic diagram of deep neck space.

head and neck, extending through the visceral space and even reaching the mediastinum [31, 32]. Second, such a large amount of liquid led only to rightward deviation of the trachea, without causing marked narrowing in the diameter or airway compression. Ultimately, the monkey woke up 1 h after the last CT examination and survived for >1 month.

The animal experiment revealed that the hydrodissection liquid did not behave like post-thyroidectomy hemorrhage, which causes serious complications, including airway compression, when the volume is approximately 50 mL [33, 34]. This occurs because postoperative hemorrhage quickly coagulates into blood clots and thus compresses the trachea. However, the injected liquid lacked blood cells and coagulation factors. The lack of coagulation components provided the hydrodissection liquid with good flowability and thus allowed it to spread freely. Not only did the liquid not agglutinate, but it also spread to the space around the thyroid. Then, the full diameter of the trachea ensured sufficient air inhalation and did not affect the vital signs or survival of the monkey. This is the first study to report the fate of hydrodissection liquid injected around the thyroid in animal experiments. The animal study theoretically demonstrated the safety of a large volume of hydrodissection liquid.

Several limitations should be acknowledged. First, the safety threshold of the hydrodissection volume has not been shown. Although no complications occurred in the 2 patients who were injected with as much as 450 mL of liquid, it is still impossible to answer whether 450 mL or more liquid is the upper limit for hydrodissection vol-

umes. Fortunately, by monitoring the amount of fluid around the thyroid and supplementing it in a timely manner, the 3- to 5-mm safety buffer was shown to be effective and safe. Second, given that this is a retrospective cohort study, there are inherent problems related to the study design. Third, there were some differences between the rhesus monkey and humans when the hydrodissection liquid was injected. The rhesus monkey did not undergo ablation, was under general anesthesia instead of local anesthesia, and changed postures during the procedure. As a result, the distribution and movement of the liquid in the animal may have differed slightly from those in patients.

Conclusion

Continuous, large-volume hydrodissection could protect the vital structures around the thyroid throughout the RFA procedure, which might be particularly beneficial for large nodule ablation. Furthermore, animal experiments showed a potentially large space around the neck, and the mediastinum was able to store a large volume of hydrodissection liquid without causing airway compression.

Statement of Ethics

This study was approved by the Review Board of the Third Affiliated Hospital of Sun Yat-sen University and was registered at www.chictr.org.cn (ChiCTR-INR-16007884). The animal study was approved by the Institutional Animal Care and Use Committee (IACUC) of Sun Yat-sen University (No. 2018-175).

Conflict of Interest Statement

All authors have no conflicts of interest to declare.

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Author Contributions

Yanping Ma contributed to literature research, experimental studies, statistical analysis, and manuscript drafting. Tao Wu contributed to clinical studies, experimental studies, and manuscript

drafting. Zhicheng Yao contributed to literature research, experimental studies, and manuscript drafting. Bowen Zheng contributed to clinical studies and statistical analysis. Lei Tan contributed to clinical studies. Ge Tong contributed to statistical analysis. Yufan Lian contributed to statistical analysis. Jung Hwan Baik contributed to study design and manuscript editing. Jie Ren contributed to manuscript editing and study design and is the guarantor of integrity of the entire study.

Data Availability Statement

The data that support the findings of this study are not publicly available due to their containing information that could compromise the privacy of research participants but are available from Ma (mayp3@mail2.sysu.edu.cn) upon reasonable request.

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